



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 176541

TO: Stephen Kapushoc
Location: 3a60 / 2c70
Tuesday, January 17, 2006
Art Unit: 1634
Phone: 571-272-3312
Serial Number: 10 / 754446

From: Jan Delaval
Location: Biotech-Chem Library
Remsen 1a51
Phone: 571-272-2504

jan.delaval@uspto.gov

Search Notes

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STIC-Biotech/ChemLib

176 513

ME

From: Kapushoc, Stephen T.
Sent: Thursday, January 12, 2006 11:15 AM
To: STIC-Biotech/ChemLib
Subject: Sequence searches for 10/754,446

RECEIVED
JAN 12 2006
STIC

Please perform the following searches for application 10/754,446:

Score/Length sequence search for the following fragments of GenBank Locus AF287270, with a minimum hit length of 15 and a maximum hit length of 30, and a score over length cutoff at 90%:
Fragments of AF287270: 100-500, 6956-7356

Please also search the following sequences: SEQ ID NOs: 1, 2, 3, 4, 5, 6, 7

441-460 = 1-20 SEQ 3

Thanks,
Steve

Stephen Kapushoc
Art Unit 1634 - USPTO
Tel: 571-272-3312
Office: REM 3A60
Mailbox: REM 2C70

09/851494

Goldin + Scierno

Searcher: Qar
Searcher Phone: 22504
Date Searcher Picked up: 1/12/06
Date completed: 1/17/06
Searcher Prep Time: 10
Online Time: (50)

Type of Search
NA# ✓ AA#
S/L: c Oligomer:
Encode/Transl:
Structure #: Text:
Inventor: Litigation:

Vendors and cost where applicable
STN:
DIALOG:
QUESTEL/ORBIT:
LEXIS/NEXIS:
SEQUENCE SYSTEM: ✓
WWW/Internet:
Other (Specify):

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 14:53:29 ; Search time 1 Seconds
(without alignments)
2.849 Million cell updates/sec

Title: af287270
Perfect score: 401
Sequence: 1 GGGGTCTCAGCAGGACAA.....TAGCTCACTGCGCCTCGAC 401

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 0.5

Searched: 189 seqs, 3552 residues

Total number of hits satisfying chosen parameters: 378

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 189 summaries

Database : af287270_copy_100_500.rng4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	25	6.2	25	1	Human mucopolip (M
2	23	5.7	24	1	Human PBA6-06 PCR
3	22.4	5.6	24	1	Human leucine zipp
4	21	5.2	21	1	Human NF-kappaB as
5	21	5.2	21	1	Human NF-kappaB as
6	20.4	5.1	22	1	Primer detects mar
7	20.4	5.1	22	1	Primer detects mar
8	20.4	5.1	22	1	Primer detects mar
9	20.4	5.1	22	1	Primer detects mar
10	20.4	5.1	22	1	Primer K to isolat
11	20.4	5.1	22	1	AHRECASEPO transge
12	20.4	5.1	22	1	Human chromosome 1
13	20	5.0	20	1	PCR primer SRI use
14	20	5.0	20	1	Primer used to amp
15	20	5.0	20	1	PCR primer SRI use
16	20	5.0	20	1	Human PDE4C oligon
17	20	5.0	20	1	Human PDE4C-deri
18	20	5.0	20	1	Human PTPN12 antis
19	20	5.0	20	1	Oligonucleotide as
20	20	5.0	20	1	Phosphorothioate a
21	20	5.0	20	1	Human oligonucleot
22	20	5.0	20	1	Single multi-plex P
23	20	5.0	20	1	Human (MCO) gene
24	20	5.0	21	1	Ataxia-telangiecta
25	20	5.0	21	1	Human AGT-105 gene
26	19.4	4.8	21	1	Primer G to isolat
27	19.4	4.8	21	1	L1 cleavage site r
28	19.4	4.8	21	1	L1 retrotransposon
29	19.4	4.8	21	1	Non-nucleotide pro
30	19	4.7	19	1	SNP specific lower
31	19	4.7	19	1	Human interleukin-
32	19	4.7	19	1	Human interleukin-
33	19	4.7	19	1	MAPK9 marker ampl

sun 1
Hokush

1	ADX86423	19	4.7	XIAP targeting sir
2	ADX86890	19	4.7	XIAP targeting sir
3	AAZ37713	20	4.7	Human mdm2 phospho
4	AAZ21805	20	4.7	Exemplary oligonuc
5	AAF80867	20	4.7	Human mdm2 phospho
6	AAZ29482	20	4.7	Human mdm2 antisen
7	ADZ21678	20	4.7	Human mdm2 antisen
8	ABZ98013	20	4.7	Human RANTES oligo
9	ABD31044	20	4.7	Human RANTES-deri
10	ADJ59878	20	4.7	Oligonucleotide as
11	ADM14235	20	4.7	Human mPGEs-1 chim
12	ADM14208	20	4.7	Human mPGEs-1 chim
13	ADO45368	20	4.7	Human oligonucleot
14	ADZ42364	20	4.7	RT-PCR primer used
15	AAZ65956	20	4.7	Primer #1 to ampli
16	AAZ37712	20	4.6	Human mdm2 phospho
17	AAZ37711	20	4.6	Human mdm2 phospho
18	AAF80866	20	4.6	Human mdm2 phospho
19	AAF80865	20	4.6	Human mdm2 phospho
20	AAH20695	20	4.6	Human telomeric re
21	AAZ29480	20	4.6	Human mdm2 antisen
22	AAZ29481	20	4.6	Human mdm2 antisen
23	AAZ12408	20	4.6	Human caspase 8 mR
24	ABS67840	20	4.6	Human casein kinas
25	ADD21676	20	4.6	Human mdm2 antisen
26	ADD21677	20	4.6	Human mdm2 antisen
27	ADD25037	20	4.6	Human caspase-8 an
28	ABZ98012	20	4.6	Human RANTES oligo
29	ABZ99087	20	4.6	Human PDE4C oligon
30	ABZ99103	20	4.6	Human PDE4C oligon
31	ABD31043	20	4.6	Human RANTES-deri
32	ABD32134	20	4.6	Human PDE4C-deri
33	ABD32118	20	4.6	Human PDE4C-deri
34	ADJ59877	20	4.6	Oligonucleotide as
35	ADJ60972	20	4.6	Oligonucleotide as
36	ADJ60988	20	4.6	Oligonucleotide as
37	ADJ96297	20	4.6	Human breast cance
38	ADJ96333	20	4.6	Human breast cance
39	ADJ96333	20	4.6	Human breast cance
40	ADJ96457	20	4.6	Human breast cance
41	ADM14854	20	4.6	Human mPGEs-1 chim
42	ADM14914	20	4.6	Human mPGEs-1 chim
43	ADM14565	20	4.6	Human mPGEs-1 chim
44	ADM15136	20	4.6	Human mPGEs-1 chim
45	ADM15380	20	4.6	Human mPGEs-1 chim
46	ADM14196	20	4.6	Human mPGEs-1 chim
47	ADM14761	20	4.6	Human mPGEs-1 chim
48	ADM15044	20	4.6	Human mPGEs-1 chim
49	ADM14980	20	4.6	Human mPGEs-1 chim
50	ADO46461	20	4.6	Human oligonucleot
51	ADO46477	20	4.6	Human oligonucleot
52	ADO45367	20	4.6	Human breast cance
53	ADW98085	20	4.6	Human breast cance
54	ADW98258	20	4.6	Human breast cance
55	ADW98121	20	4.6	Human breast cance
56	ADW98194	20	4.6	Human breast cance
57	ADX82227	20	4.6	Melanoma associate
58	ADX82226	20	4.6	Melanoma associate
59	ADY92906	20	4.6	PCR primer U479 us
60	AAQ25869	19	4.5	3' Alu primer. Sy
61	AAQ48683	19	4.5	Human Alu segment
62	AAQ85677	19	4.5	PCR primer alu 2 f
63	ADZ74318	19	4.5	Inter-Alu PCR prim
64	ADY01152	18	4.5	Extend primer 28 u
65	AAZ01233	19	4.5	Forward PCR primer
66	ADH99039	19	4.5	Human POLYX PCR pr
67	ADY03156	19	4.5	Extend primer 606
68	AAZ61524	20	4.5	Human inhibitor-ka
69	ABX93650	19	4.4	Human Alu-specific
70	ABX95026	19	4.4	Human Alu specific
71	AAH39033	19	4.3	SNP specific upper
72	AAH24568	19	4.3	Human Alu sequence

c 107	17.4	4.3	19	1	ADY01278	Extend primer 154					
c 108	17.4	4.3	19	1	ADX86792	XIAP targeting sir					
c 109	17.4	4.3	19	1	ADX86325	XIAP targeting sir					
c 110	17.4	4.3	19	1	ABE05189	Human IL-13RA1 sir					
c 111	17.4	4.3	19	1	ABE05166	Human IL-13RA1 tra					
c 112	17.4	4.3	19	1	AEC15226	Human IL-13RA1 sir					
c 113	17.4	4.3	19	1	AEC15003	Human IL-13RA1 tra					
c 114	17.2	4.3	19	1	AAQ76248	Generic primer fro					
c 115	17	4.2	17	1	AAV23284	Nucleotide sequence					
c 116	17	4.2	17	1	AAC87597	Human Alu sequence					
c 117	17	4.2	17	1	ADB44570	Tumour suppression					
c 118	17	4.2	17	1	ACC51496	Human tumour suppr					
c 119	17	4.2	17	1	ADL50193	Human PKR substrat					
c 120	17	4.2	17	1	ADL50731	Human PKR substrat					
c 121	17	4.2	17	1	ADL49907	Human PKR substrat					
c 122	17	4.2	17	1	ADL49908	Human PKR substrat					
c 123	17	4.2	17	1	ADL49909	Human PKR substrat					
c 124	17	4.2	17	1	ADL49906	Human PKR substrat					
c 125	17	4.2	17	1	ADL50192	Human PKR substrat					
c 126	17	4.2	17	1	ADL82338	Human ER+ breast c					
c 127	17	4.2	17	1	ADP08690	Extend primer 27 u					
c 128	17	4.2	17	1	ADY01178	Extend primer 54 u					
c 129	17	4.2	17	1	ADY00434	Extend primer 2 us					
c 130	17	4.2	17	1	ADX82001	Melanoma associate					
c 131	17	4.2	17	1	ADX81510	Melanoma associate					
c 132	17	4.2	18	1	ADO56979	Human CARK/FPGT pr					
c 133	17	4.2	18	1	ADO56561	Human cyclin-depen					
c 134	16.4	4.1	18	1	AAZ27769	PCR primer for hum					
c 135	16.4	4.1	18	1	ADY02910	Extend primer 360					
c 136	16.4	4.1	18	1	ADY00671	Extend primer 239					
c 137	16.4	4.1	18	1	ADY01277	Extend primer 153					
c 138	16	4.0	16	1	AAF88161	Human thyroid malf					
c 139	16	4.0	16	1	ACA62885	Repeated nucleic a					
c 140	16	4.0	16	1	ACA62882	Repeated nucleic a					
c 141	16	4.0	17	1	ADB04313	Human MDZ7 scannin					
c 142	16	4.0	17	1	ADB04314	Human MDZ7 scannin					
c 143	16	4.0	17	1	ACC63031	Murine oligonucleo					
c 144	16	4.0	17	1	ADB44260	Tumour suppression					
c 145	16	4.0	17	1	ACC51495	Human tumour suppr					
c 146	16	4.0	17	1	ADL50732	Human PKR substrat					
c 147	16	4.0	17	1	ADL49420	Human PKR substrat					
c 148	16	4.0	17	1	ADX81033	Melanoma associate					
c 149	15.4	3.8	17	1	AAA22718	Integrin subunit b					
c 150	15.4	3.8	17	1	AAA22717	Integrin subunit b					
c 151	15.4	3.8	17	1	ABT39409	Tumour suppression					
c 152	15.4	3.8	17	1	ABT38213	Tumour suppression					
c 153	15.4	3.8	17	1	ABT38720	Tumour suppression					
c 154	15.4	3.8	17	1	ABT38728	Tumour suppression					
c 155	15.4	3.8	17	1	ABT35457	Tumour suppression					
c 156	15.4	3.8	17	1	ABT40068	Tumour suppression					
c 157	15.4	3.8	17	1	ABT40150	Tumour suppression					
c 158	15.4	3.8	17	1	ABT35874	Tumour suppression					
c 159	15.4	3.8	17	1	ABZ60567	Human K-Ras DNazym					
c 160	15.4	3.8	17	1	ADB41143	Tumour suppression					
c 161	15.4	3.8	17	1	ADB43523	Tumour suppression					
c 162	15.4	3.8	17	1	ADB44518	Tumour suppression					
c 163	15.4	3.8	17	1	ADE43565	Human IDE sequenci					
c 164	15.4	3.8	17	1	ADI50915	Human tumour suppr					
c 165	15.4	3.8	17	1	ADI50723	Human tumour suppr					
c 166	15.4	3.8	17	1	ADI52180	Human tumour suppr					
c 167	15.4	3.8	17	1	ADI50051	Human tumour suppr					
c 168	15.4	3.8	17	1	ADI51643	Human tumour suppr					
c 169	15.4	3.8	17	1	ACC52610	Human tumour suppr					
c 170	15.4	3.8	17	1	ACC51497	Human tumour suppr					
c 171	15.4	3.8	17	1	ACC52025	Human tumour suppr					
c 172	15.4	3.8	17	1	ACC53324	Human tumour suppr					
c 173	15.4	3.8	17	1	ADL49905	Human PKR substrat					
c 174	15.4	3.8	17	1	ADL49419	Human PKR substrat					
c 175	15.4	3.8	17	1	ADL49904	Human PKR substrat					
c 176	15.4	3.8	17	1	ADL50191	Human PKR substrat					
c 177	15.4	3.8	17	1	ADL50733	Human PKR substrat					
c 178	15.4	3.8	17	1	ADH54043	Human neurodegener					
c 179	15.4	3.8	17	1	ADK13186	Human glioma endot					
c 180	15.4	3.8	17	1	ADL82349	Human ER+ breast c					
c 181	15.4	3.8	17	1	ADZ29641	Human K-Ras subestr					
c 182	15.4	3.8	17	1	AEF71294	Extension primer f					
c 183	15	3.7	15	1	AAF69438	Human IL4Ralpha ge					
c 184	14	3.5	15	1	ABK98169	Triple helix formi					
c 185	14	3.5	15	1	ABK98187	Triple helix formi					
c 186	14	3.5	15	1	ABK98168	Triple helix formi					
c 187	14	3.5	15	1	ABK98167	Triple helix formi					
c 188	14	3.5	15	1	ABK98186	Triple helix formi					
c 189	13.6	3.4	15	1	AAS98701	Colony stimulating					
ALIGNMENTS											
RESULT 1											
ADB84281											
ID	ADB84281 standard; DNA; 25 BP.										
XX	AC ADB84281;										
XX	04-DEC-2003 (first entry)										
XX	Human mucolipin (MCOLIPIN-1) gene AJ haplotype analysis primer #3.										
XX	human; mucolipin; MCOLN-1; mucopolipidosis IV; ion channel defect;										
KW	immunogen; gene therapy; ss; primer; PCR; AJ haplotype.										
XX	Homo sapiens.										
XX	US2003064363-A1.										
XX	03-APR-2003.										
PF	08-MAY-2001; 2001US-00851494.										
XX	18-AUG-2000; 2000US-0226388P.										
XX	(MLFO-) ML4 FOUND & HAVARD COLLEGE.										
XX	Goldin E, Slaugenhaupt SA, Sun M, Acierno JS;										
XX	WPI; 2003-540779/51.										
XX	New mucolipin (MCOLN1) polypeptides and nucleic acids, useful in gene										
PT	therapy for treating mucopolipidosis IV or ion channel defect, or as										
PT	immunogen to generate antibodies that recognize the MCOLN1 polypeptide.										
XX	Example; Page 18; 34pp; English.										
CC	The invention describes a nucleic acid (I) encoding a mucolipin (MCOLN1)										
CC	polypeptide, and sharing at least 95% sequence identity with sequence										
CC	selected from 2 nucleotide sequences (designated S1 and S2; not given in										
CC	the specification). A mutation of in the MCOLN1 gene encoding the MCOLN1										
CC	polypeptide results in a defect in expression of a functional MCOLN1. The										
CC	MCOLN1 gene is useful for the treatment of mucopolipidosis IV or										
CC	ion channel defect. The MCOLN1 polypeptides are useful as immunogen to										
CC	generate antibodies that recognize the MCOLN1 polypeptide. Such										
CC	antibodies may be used for diagnostics and intracellular regulation of										
CC	MCOLN1 activity. The polynucleotides may also be useful in gene therapy.										
CC	This sequence represents a primer used to analyse the AJ haplotype of the										
CC	human mucolipin (MCOLN-1) gene.										
XX	SQ Sequence 25 BP; 7 A; 5 C; 6 G; 7 T; 0 U; 0 Other;										
Query Match 6.2%; Score 25; DB 1; Length 25;											
Best Local Similarity 100.0%; Pred. No. 12;											
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;											
QY	226 CTGATATAAATGGCAGGCAGCTTTC 250										
DB	1 CTGATATAAATGGCAGGCAGCTTTC 25										

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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 14:56:42 ; Search time 2 Seconds
(without alignments)
2.069 Million cell updates/sec

Title: af287270

Perfect score: 401

Sequence: 1 GGGTCTCAGCAGGGAACAA.....TAGCTCAGCGCTCGAC 401

Scoring table: IDENTITY NUC

Gap 10.0 , Gapext 0.5

Searched: 244 seqs, 5160 residues

Total number of hits satisfying chosen parameters: 488

Minimum DB seq length: 15

Maximum DB seq length: 30

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 244 summaries

Database : af287270_copy_100_500.rnpbm4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	25.4	6.3	27	1 US-10-708-204-5106	Sequence 5106, Ap
2	25	6.2	25	1 US-10-956-157-173066	Sequence 173066, Ap
3	25	6.2	25	1 US-11-060-756-178198	Sequence 178198, Ap
4	24	6.0	24	1 US-10-708-204-5162	Sequence 5162, Ap
5	24	6.0	25	1 US-10-956-157-185762	Sequence 185762, Ap
6	24	6.0	25	1 US-10-956-157-208929	Sequence 208929, Ap
7	24	6.0	25	1 US-11-060-756-178197	Sequence 178197, Ap
8	24	6.0	25	1 US-11-060-756-203216	Sequence 203216, Ap
9	23.4	5.8	25	1 US-10-956-157-149516	Sequence 149516, Ap
10	23.4	5.8	25	1 US-10-956-157-149517	Sequence 149517, Ap
11	23.4	5.8	25	1 US-10-956-157-173133	Sequence 173133, Ap
12	23.4	5.8	25	1 US-10-956-157-177068	Sequence 177068, Ap
13	23.4	5.8	25	1 US-10-956-157-177069	Sequence 177069, Ap
14	23.4	5.8	25	1 US-10-956-157-184259	Sequence 184259, Ap
15	23.4	5.8	25	1 US-10-956-157-188388	Sequence 188388, Ap
16	23.4	5.8	25	1 US-10-956-157-273920	Sequence 273920, Ap
17	23.4	5.8	25	1 US-10-956-157-285780	Sequence 285780, Ap
18	23.4	5.8	25	1 US-10-956-157-286030	Sequence 286030, Ap
19	23.4	5.8	25	1 US-10-956-157-287740	Sequence 287740, Ap
20	23.4	5.8	25	1 US-10-956-157-293223	Sequence 293223, Ap
21	23.4	5.8	25	1 US-10-956-157-293224	Sequence 293224, Ap
22	23.4	5.8	25	1 US-10-956-157-293225	Sequence 293225, Ap
23	23.4	5.8	25	1 US-10-956-157-298488	Sequence 298488, Ap
24	23.4	5.8	25	1 US-11-060-756-5164	Sequence 5164, Ap
25	23.4	5.8	25	1 US-11-060-756-163912	Sequence 163912, Ap
26	23.4	5.8	25	1 US-11-060-756-164006	Sequence 164006, Ap
27	23.4	5.8	25	1 US-11-060-756-180494	Sequence 180494, Ap
28	23.4	5.8	25	1 US-11-060-756-200858	Sequence 200858, Ap
29	23.4	5.8	25	1 US-11-060-756-272590	Sequence 272590, Ap
30	23.4	5.8	25	1 US-11-060-756-274785	Sequence 274785, Ap
31	23.4	5.8	25	1 US-11-060-756-278943	Sequence 278943, Ap
32	23.4	5.8	25	1 US-11-060-756-279896	Sequence 279896, Ap
33	23.4	5.8	25	1 US-11-060-756-279897	Sequence 279897, Ap

25	5.8	23.4	1	US-11-060-756-287805	Sequence 287805, Ap
23	5.7	23	1	US-09-851-494B-10	Sequence 10, Appl
23	5.7	23	1	US-10-708-204-5076	Sequence 5076, Ap
23	5.7	23	1	US-10-708-204-5099	Sequence 5099, Ap
22	5.5	22	1	US-10-708-204-7	Sequence 7, Appl
22	5.5	22	1	US-10-708-204-13	Sequence 13, Appl
22	5.5	22	1	US-10-708-204-72	Sequence 72, Appl
22	5.5	22	1	US-10-708-204-91	Sequence 91, Appl
22	5.5	22	1	US-10-708-204-195	Sequence 195, Appl
22	5.5	22	1	US-10-708-204-208	Sequence 208, Appl
22	5.5	22	1	US-10-708-204-771	Sequence 771, Appl
22	5.5	22	1	US-10-708-204-1122	Sequence 1122, Appl
22	5.5	22	1	US-10-708-204-1127	Sequence 1127, Appl
22	5.5	22	1	US-10-708-204-1177	Sequence 1177, Appl
22	5.5	22	1	US-10-708-204-1399	Sequence 1399, Appl
22	5.5	22	1	US-10-708-204-1401	Sequence 1401, Appl
22	5.5	22	1	US-10-708-204-1480	Sequence 1480, Appl
22	5.5	22	1	US-10-708-204-1498	Sequence 1498, Appl
22	5.5	22	1	US-10-708-204-2082	Sequence 2082, Appl
22	5.5	22	1	US-10-708-204-2092	Sequence 2092, Appl
22	5.5	22	1	US-10-708-204-2517	Sequence 2517, Appl
22	5.5	22	1	US-10-708-204-2540	Sequence 2540, Appl
22	5.5	22	1	US-10-708-204-301	Sequence 301, Appl
22	5.5	22	1	US-10-708-204-4301	Sequence 4301, Appl
22	5.5	22	1	US-10-708-204-5075	Sequence 5075, Appl
22	5.5	22	1	US-10-708-204-5086	Sequence 5086, Appl
22	5.5	22	1	US-10-708-204-5095	Sequence 5095, Appl
22	5.5	22	1	US-10-708-204-5098	Sequence 5098, Appl
22	5.5	22	1	US-10-708-204-5105	Sequence 5105, Appl
22	5.5	22	1	US-10-708-204-3821	Sequence 3821, Appl
22	5.5	22	1	US-10-708-204-3863	Sequence 3863, Appl
22	5.5	22	1	US-10-708-204-5124	Sequence 5124, Appl
22	5.5	22	1	US-10-126-103-235	Sequence 235, Appl
22	5.5	22	1	US-10-431-096-235	Sequence 235, Appl
22	5.5	22	1	US-10-708-204-3851	Sequence 3851, Appl
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22	5.5	22	1	US-10-708-204-5096	Sequence 5096, Appl
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22	5.5	22	1	US-10-708-204-5097	Sequence 5097, Appl
22	5.5	22	1	US-09-918-686-93	Sequence 93, Appl
22	5.5	22	1	US-10-353-150-93	Sequence 93, Appl
22	5.5	22	1	US-10-374-077-11	Sequence 11, Appl
22	5.5	22	1	US-10-708-204-14	Sequence 14, Appl
22	5.5	22	1	US-10-708-204-73	Sequence 73, Appl
22	5.5	22	1	US-10-708-204-75	Sequence 75, Appl
22	5.5	22	1	US-10-708-204-162	Sequence 162, Appl
22	5.5	22	1	US-10-708-204-192	Sequence 192, Appl
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c 113	20.4	5.1	22	1	US-10-708-204-2248	Sequence 2248, Ap	c 186	18.4	4.6	20	1	US-10-671-395-1167	Sequence 1167, Ap
c 114	20.4	5.1	22	1	US-10-708-204-2249	Sequence 2249, Ap	c 187	18.4	4.6	20	1	US-10-671-395-1231	Sequence 1231, Ap
c 115	20.4	5.1	22	1	US-10-708-204-2250	Sequence 2250, Ap	c 188	18.4	4.6	20	1	US-10-671-395-1323	Sequence 1323, Ap
c 116	20.4	5.1	22	1	US-10-708-204-2252	Sequence 2252, Ap	c 189	18.4	4.6	20	1	US-10-671-395-1549	Sequence 1549, Ap
117	20.4	5.1	22	1	US-10-708-204-2516	Sequence 2516, Ap	c 190	18.4	4.6	20	1	US-10-671-395-1567	Sequence 1567, Ap
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119	20.4	5.1	22	1	US-10-708-204-3820	Sequence 3820, Ap	c 192	18.4	4.6	20	1	US-10-639-300-74	Sequence 74, Appl
120	20.4	5.1	22	1	US-10-708-204-3827	Sequence 3827, Ap	c 193	18.4	4.6	20	1	US-10-840-590-1509	Sequence 1509, Ap
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c 124	20	5.0	20	1	US-10-251-699-1	Sequence 1, Appli	c 197	18	4.5	19	1	US-10-098-871-37	Sequence 37, Appl
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c 136	20	5.0	22	1	US-10-708-204-831	Sequence 831, App	c 209	17.4	4.3	19	1	US-11-014-373-467	Sequence 467, App
c 137	20	5.0	22	1	US-10-708-204-876	Sequence 876, App	c 210	17.4	4.3	19	1	US-10-463-9818-2	Sequence 2, Appli
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c 140	20	5.0	22	1	US-10-708-204-1102	Sequence 1102, Ap	c 213	17	4.2	17	1	US-10-156-306-1652	Sequence 1652, Ap
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143	20	5.0	22	1	US-10-708-204-1487	Sequence 1487, Ap	c 216	17	4.2	17	1	US-10-156-306-2389	Sequence 2389, Ap
c 144	20	5.0	22	1	US-10-708-204-1539	Sequence 1539, Ap	c 217	17	4.2	17	1	US-10-156-306-2390	Sequence 2390, Ap
c 145	20	5.0	22	1	US-10-708-204-1653	Sequence 1653, Ap	c 218	17	4.2	17	1	US-10-156-306-3776	Sequence 3776, Ap
c 146	20	5.0	22	1	US-10-708-204-2243	Sequence 2243, Ap	c 219	17	4.2	17	1	US-10-339-782-309	Sequence 309, App
c 147	20	5.0	22	1	US-10-708-204-2274	Sequence 2274, Ap	c 220	17	4.2	17	1	US-10-704-513-302	Sequence 302, App
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c 149	20	5.0	22	1	US-10-708-204-4312	Sequence 4312, Ap	c 222	17	4.2	17	1	US-10-840-590-793	Sequence 793, App
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c 151	20	5.0	22	1	US-10-708-204-4385	Sequence 4385, Ap	c 224	17	4.2	17	1	US-09-784-423-69	Sequence 69, Appl
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c 155	19.4	4.8	21	1	US-10-255-434-25	Sequence 25, Appl	c 228	16	4.0	16	1	US-10-156-306-534	Sequence 534, App
c 156	19.4	4.8	21	1	US-10-374-077-7	Sequence 7, Appli	c 229	16	4.0	17	1	US-10-156-306-3777	Sequence 3777, Ap
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158	19	4.7	19	1	US-10-708-204-3840	Sequence 3840, Ap	c 231	16	4.0	17	1	US-10-840-590-314	Sequence 314, App
159	19	4.7	19	1	US-10-708-204-3842	Sequence 3842, Ap	c 232	16	4.0	17	1	US-10-156-306-533	Sequence 533, App
160	19	4.7	19	1	US-10-708-204-3854	Sequence 3854, Ap	c 233	15.4	3.8	17	1	US-10-156-306-1649	Sequence 1649, Ap
161	19	4.7	19	1	US-10-708-204-4379	Sequence 4379, Ap	c 234	15.4	3.8	17	1	US-10-156-306-1650	Sequence 1650, Ap
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c 163	19	4.7	19	1	US-10-708-204-5172	Sequence 5172, Ap	c 236	15.4	3.8	17	1	US-10-156-306-3778	Sequence 3778, Ap
c 164	19	4.7	20	1	US-09-752-983-243	Sequence 243, App	c 237	15.4	3.8	17	1	US-10-339-782-320	Sequence 320, App
c 165	19	4.7	20	1	US-09-863-806-155	Sequence 155, App	c 238	15.4	3.8	17	1	US-10-339-793-16	Sequence 16, Appl
c 166	19	4.7	20	1	US-10-005-344-243	Sequence 243, App	c 239	15.4	3.8	17	1	US-10-600-009-170	Sequence 170, App
c 167	19	4.7	20	1	US-10-671-395-395	Sequence 395, App	c 240	15.4	3.8	17	1	US-10-724-270-679	Sequence 679, App
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c 169	19	4.7	20	1	US-10-754-478-155	Sequence 155, App	c 242	15.4	3.8	17	1		
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171	19	4.7	21	1	US-10-708-204-4384	Sequence 4384, Ap	c 244	15	3.7	15	1		
172	19	4.7	21	1	US-10-708-204-5102	Sequence 5102, Ap							
c 173	18.4	4.6	20	1	US-09-752-983-241	Sequence 241, App							
c 174	18.4	4.6	20	1	US-09-752-983-242	Sequence 242, App							
c 175	18.4	4.6	20	1	US-10-181-177-94	Sequence 94, Appl							
c 176	18.4	4.6	20	1	US-10-005-344-241	Sequence 241, App							
c 177	18.4	4.6	20	1	US-10-005-344-242	Sequence 242, App							
c 178	18.4	4.6	20	1	US-10-148-355A-64	Sequence 64, Appl							
179	18.4	4.6	20	1	US-10-199-676-38	Sequence 38, Appl							

RESULT 1
US-10-708-204-5106
; Sequence 5106, Application us/10708204

ALIGNMENTS

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 14:58:13 ; Search time 0.001 Seconds
(without alignments)
318.394 Million cell updates/sec

Title: af287270
Perfect score: 401
Sequence: 1 GGGTCTCAGCAGGGAACAA.....TAGTCACTGAGCCTCGAC 401

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 0.5

Searched: 19 segs, 397 residues

Total number of hits satisfying chosen parameters: 38

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 19 summaries

Database : af287270_copy_100_500.rnpbn4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	23.4	5.8	25	1	US-11-121-849-44508
C 3	23.4	5.8	25	1	US-11-121-849-44509
C 4	23.4	5.8	25	1	US-11-121-849-48982
C 5	23.4	5.8	25	1	US-11-121-849-48983
C 6	23.4	5.8	25	1	US-11-121-849-48984
C 7	23.4	5.8	25	1	US-11-121-849-48985
C 8	20	5.0	20	1	US-10-515-538-80
C 9	19	4.7	20	1	US-10-353-783-33
C 10	18	4.5	18	1	US-10-857-780-2580
C 11	18	4.5	19	1	US-10-857-780-4587
C 12	17.4	4.3	19	1	US-10-857-780-2706
C 13	17.4	4.3	19	1	US-11-001-347-922
C 14	17.4	4.3	19	1	US-11-001-347-1145
C 15	17	4.2	17	1	US-10-857-780-1868
C 16	17	4.2	17	1	US-10-857-780-2606
C 17	16.4	4.1	18	1	US-10-857-780-2105
C 18	16.4	4.1	18	1	US-10-857-780-2705
C 19	16.4	4.1	18	1	US-10-857-780-4341

ALIGNMENTS

RESULT 1
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; Sequence 17198, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE OF INVENTION: Microarrays
; FILE REFERENCE: 3684.1

; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 17198
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-17198

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Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 25 TCTTGCTCTGTTGCCAGGCTGGAG 1

RESULT 2

US-11-121-849-44508/c
; Sequence 44508, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE OF INVENTION: Microarrays
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 44508
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-44508

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Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 3

US-11-121-849-44509/c
; Sequence 44509, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE OF INVENTION: Microarrays
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 44509
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-44509

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Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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RESULT 4
US-11-121-849-48982/c
; Sequence 48982, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 48982
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-48982

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Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 25 CTGTGCCCCAGGCTGGAGTGCAGTG 1

RESULT 5
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; Sequence 48983, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 48983
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-48983

Query Match 5.8%; Score 23.4; DB 1; Length 25;
Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 446 TCTGTGCCCCAGGCTGGAGTGCACAT 470
Db 25 TCTGTGCCCCAGGCTGGAGTGCACAT 1

RESULT 6
US-11-121-849-48984/c
; Sequence 48984, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
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; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 48984
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-48984

Query Match 5.8%; Score 23.4; DB 1; Length 25;
Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 445 CTCTGTGCCCCAGGCTGGAGTGCAG 469
Db 25 CTCTGTGCCCCAGGCTGGAGTGCAG 1

RESULT 7
US-11-121-849-48985/c
; Sequence 48985, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR APPLICATION NUMBER: 60/567,949
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 48985
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-48985

Query Match 5.8%; Score 23.4; DB 1; Length 25;
Best Local Similarity 96.0%; Pred. No. 2;
Matches 24; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 444 GCTCTGTGCCCCAGGCTGGAGTGCA 468
Db 25 GCTCTGTGCCCCAGGCTGGAGTGCA 1

RESULT 8
US-10-515-538-80
; Sequence 80, Application US/10515538
; Publication No. US20050282760A1
; GENERAL INFORMATION:
; APPLICANT: Isis Pharmaceuticals, Inc.
; APPLICANT: Lex M. Cowsett
; APPLICANT: Kenneth W. Dobie
; TITLE OF INVENTION: ANTISENSE MODULATION OF PTPN12 EXPRESSION
; FILE REFERENCE: PFS-0016USA
; CURRENT APPLICATION NUMBER: US/10/515,538
; CURRENT FILING DATE: 2004-11-23
; PRIOR APPLICATION NUMBER: 10/172,911
; PRIOR FILING DATE: 2002-06-17
; PRIOR APPLICATION NUMBER: PCT/US03/18707
; PRIOR FILING DATE: 2003-06-12
; NUMBER OF SEQ ID NOS: 123
; SEQ ID NO 80
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 14:54:56 ; Search time 0.001 Seconds
(without alignments)
874.180 Million cell updates/sec

Title: af287270

Perfect score: 401

Sequence: 1 GGGGTCTCAGCAGGACAA.....TAGCTCACTGAGCCTCGAC 401

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 0.5

Searched: 58 seqs, 1090 residues

Total number of hits satisfying chosen parameters: 116

Minimum DB seq length: 15

Maximum DB seq length: 30

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 58 summaries

Database : af287270_copy_100_500.rn14:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	20.4	5.1	22	1	US-08-599-252-35
C 2	20.4	5.1	22	1	US-08-599-252-38
C 3	20.4	5.1	22	1	US-08-291-074-2
C 4	20.4	5.1	22	1	US-08-781-891-11
C 5	20.4	5.1	22	1	US-09-918-686-93
C 6	20.4	5.1	22	1	US-09-618-166-11
C 7	20.4	5.1	22	1	PCT-US96-06352-35
C 8	20.4	5.1	22	1	PCT-US96-06352-38
C 9	20.4	5.1	22	1	PCT-US96-06583-35
C 10	20.4	5.1	22	1	PCT-US96-06583-38
C 11	20.4	5.1	22	1	US-09-418-804-1
C 12	20.4	5.0	20	1	US-10-172-911-80
C 13	20.4	5.0	20	1	US-10-251-699-1
C 14	20.4	5.0	21	1	US-08-753-147-28
C 15	19.4	4.8	21	1	US-08-781-891-7
C 16	19.4	4.8	21	1	US-08-847-844A-116
C 17	19.4	4.8	21	1	US-09-618-166-7
C 18	19.4	4.7	20	1	US-09-280-805-243
C 19	19.4	4.7	20	1	US-09-038-637-155
C 20	19.4	4.7	20	1	US-09-224-683-33
C 21	19.4	4.7	20	1	US-09-604-325A-33
C 22	18.4	4.6	20	1	US-08-222-177A-274
C 23	18.4	4.6	20	1	US-09-280-805-241
C 24	18.4	4.6	20	1	US-09-280-805-242
C 25	18.4	4.6	20	1	US-09-487-445-94
C 26	18.4	4.6	20	1	US-09-467-642-64
C 27	18.4	4.6	20	1	US-09-780-173A-18
C 28	18.2	4.5	19	1	US-08-070-517-2
C 29	18.2	4.5	19	1	US-08-118-441-2
C 30	18.2	4.5	19	1	US-08-422-699A-14
C 31	18.2	4.5	19	1	US-08-422-706B-14
C 32	18.2	4.5	19	1	US-08-338-579A-2
C 33	18.2	4.5	19	1	PCT-US94-09851-2

34	17.4	4.3	19	1	US-09-366-840-2	Sequence 2, Appli
35	17.2	4.3	19	1	US-10-176-884-44	Sequence 44, Appli
C 36	17	4.2	17	1	US-08-635-820A-2	Sequence 2, Appli
C 37	17	4.2	17	1	US-09-100-104-2	Sequence 2, Appli
38	17	4.2	17	1	US-10-054-295-132	Sequence 132, App
39	17	4.2	17	1	US-09-438-486A-132	Sequence 132, App
40	17	4.2	18	1	US-09-979-275A-7	Sequence 7, Appli
C 41	16.4	4.1	18	1	US-09-018-584A-69	Sequence 69, Appli
C 42	16.4	4.1	18	1	US-09-784-423-69	Sequence 69, Appli
C 43	16.2	4.0	18	1	US-09-981-397A-1	Sequence 1, Appli
44	16	4.0	16	1	US-10-203-780-9	Sequence 9, Appli
C 45	16	4.0	16	1	US-10-054-295-131	Sequence 131, App
C 46	16	4.0	16	1	US-09-438-486A-131	Sequence 131, App
47	16	4.0	16	1	US-10-227-001-21	Sequence 21, Appli
48	16	4.0	16	1	US-10-072-975-9	Sequence 9, Appli
49	16	4.0	17	1	US-09-730-559B-107	Sequence 107, App
50	15	3.7	15	1	US-09-850-982B-4	Sequence 4, Appli
51	15	3.7	15	1	US-09-793-146-54	Sequence 54, Appli
52	15	3.7	15	1	US-09-793-146-55	Sequence 55, Appli
53	15	3.7	15	1	US-10-227-001-23	Sequence 23, Appli
54	15	3.7	15	1	US-10-384-450A-5	Sequence 5, Appli
55	15	3.7	15	1	US-10-072-975-10	Sequence 10, Appli
56	15	3.7	16	1	US-10-227-001-20	Sequence 20, Appli
57	14.4	3.6	16	1	US-09-479-005A-260	Sequence 260, App
58	14	3.5	15	1	US-10-227-001-24	Sequence 24, Appli

ALIGNMENTS

RESULT 1
US-08-599-252-35/c
; Sequence 35, Application US/08599252
; Patent No. 5705343
; GENERAL INFORMATION:
; APPLICANT: DRAYNA, DENNIS T.
; APPLICANT: FEDER, JOHN N.
; APPLICANT: GRIKKE, ANDREAS
; APPLICANT: KIMMEL, BRUCE E.
; APPLICANT: THOMAS, WINSTON J.
; APPLICANT: WOLFF, ROGER K.
; TITLE OF INVENTION: METHOD TO DIAGNOSE HEREDITARY
; TITLE OF INVENTION: HEMOCHROMATOSIS
; NUMBER OF SEQUENCES: 124
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 2000 Pennsylvania Ave. N.W., Suite 5500
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20006-1888
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/599,252
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: MURASHIGE, KATE H.
; REGISTRATION NUMBER: 29,959
; REFERENCE/DOCKET NUMBER: 9053-0001.21
; TELEPHONE: (202) 887-1500
; TELEFAX: (202) 887-0763
; TELEX: 90-4030
; INFORMATION FOR SEQ ID NO: 35:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single

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; TOPOLOGY: linear
US-08-599-252-35

Query Match          5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 10;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      444 GCTCTGTTGCCAGCGTGGAGT 465
Db      22 GCTCTATTGCCAGCGTGGAGT 1

RESULT 2
US-08-599-252-38/c
; Sequence 38, Application US/08599252
; Patent No. 5705343
; GENERAL INFORMATION:
; APPLICANT: DRAYNA, DENNIS T.
; APPLICANT: FEDER, JOHN N.
; APPLICANT: GNIERKE, ANDREAS
; APPLICANT: KIMMEL, BRUCE E.
; APPLICANT: THOMAS, WINSTON J.
; APPLICANT: WOLFF, ROGER K.
; TITLE OF INVENTION: METHOD TO DIAGNOSE HEREDITARY
; TITLE OF INVENTION: HEMOCHROMATOSIS
; NUMBER OF SEQUENCES: 124
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 2000 Pennsylvania Ave. N.W., Suite 5500
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20006-1888
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/599,252
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: MURASHIGE, KATE H.
; REGISTRATION NUMBER: 29,959
; REFERENCE/DOCKET NUMBER: 9053-0001.21
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 887-1500
; TELEFAX: (202) 887-0763
; TELEX: 90-4030
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-599-252-38

Query Match          5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 10;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      444 GCTCTGTTGCCAGCGTGGAGT 465
Db      22 GCTCTATTGCCAGCGTGGAGT 1

RESULT 3
US-08-291-074-2
; Sequence 2, Application US/08291074
; Patent No. 5959171
; GENERAL INFORMATION:
; APPLICANT: Hyttinen, Juha-Matti
; APPLICANT: Korhonen, Veli-Pekka
; APPLICANT: Janne, Juhani
; TITLE OF INVENTION: METHOD FOR THE PRODUCTION OF
; TITLE OF INVENTION: BIOLOGICALLY ACTIVE POLYPEPTIDES IN A MAMMAL'S MILK AS
; TITLE OF INVENTION: FUSION PROTEINS THAT ARE LESS ACTIVE THAN THE FREE
; TITLE OF INVENTION: POLYPEPTIDES, OR NON-ACTIVE
; NUMBER OF SEQUENCES: 6
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Adduci, Mastriani, Schaumburg & Schill
; STREET: 1140 Connecticut Avenue, N.W.
; CITY: Washington
; STATE: DC
; COUNTRY: U.S.A.
; ZIP: 20036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/291,074
; FILING DATE: 17-AUG-1994
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Kubovcik, Ronald J.
; REGISTRATION NUMBER: 25,401
; REFERENCE/DOCKET NUMBER: TUR-017
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-467-6300
; TELEFAX: 202-466-2006
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-291-074-2

Query Match          5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 10;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      464 GTGCAGTGGTGTGATCATGCT 485
Db      1 GTGCAGTGGTGTGATCATGCT 22

RESULT 4
US-08-781-891-11
; Sequence 11, Application US/08781891
; Patent No. 6090620
; GENERAL INFORMATION:
; APPLICANT: Fu, Ying-Hui
; APPLICANT: Yu, Chang-En
; APPLICANT: Oshima, Junko
; APPLICANT: Mulligan, John T.
; APPLICANT: Schellenberg, Gerald D.
; TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
; TITLE OF INVENTION: WERNER'S SYNDROME
; NUMBER OF SEQUENCES: 209
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 15:00:13 ; Search time 0.001 Seconds
(without alignments)
13.634 Million cell updates/sec

Title: af287270
Perfect score: 401
Sequence: 1 CAGTGGGGGATCCCATCA.....CATCCTTGGCCCTTACCCGCT 401

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 0.5

Searched: 1 seqs, 17 residues

Total number of hits satisfying chosen parameters: 2

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 1 summaries

Database : af287270_copy_6956_7356.rge4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	15.4	3.8	17	1 AX757826	ACCESSION:AX757826

ALIGNMENTS

RESULT 1
AX757826/c
LOCUS AX757826 17 bp DNA linear PAT 25-JUN-2003
DEFINITION Sequence 1147 from Patent WO03040369.
ACCESSION AX757826
VERSION AX757826.1 GI:32252442
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1
AUTHORS Telerman,A., Amson,R. and Tulinder,M.
TITLE Sequences involved in tumoral suppression, tumoral reversion,
apoptosis and/or viral resistance phenomena and their use as
medicines
JOURNAL Patent: WO 03040369-A 1147 15-MAY-2003;
FEATURES Molecular Engines Laboratories (FR)
source Location/Qualifiers
1..17
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

Query Match 3.8%; Score 15.4; DB 1; Length 17;
Best Local Similarity 94.1%; Pred. No. 0;

Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 7175 CGTCCTGTGCTGAGATC 7191
||||| |||||||
Db 17 CGTCCTGGGCTGAGATC 1

Search completed: January 17, 2006, 15:00:13
Job time : 0.001 secs

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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 15:01:31 ; Search time 0.001 Seconds
(without alignments)
112.280 Million cell updates/sec

Title: af287270
Perfect score: 401
Sequence: 1 CAGTGGGGATCCCATCA.....CATCTTGGCCCTACCGCT 401

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 0.5

Searched: 8 seqs, 140 residues

Total number of hits satisfying chosen parameters: 16

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 8 summaries

Database : af287270_copy_6956_7356.rng4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	5.2	21	1	ADU28709 ✓ Knock-down target
C 2	21	5.2	21	1	ABE28960 Human MCOLN1 gene
C 3	20	5.0	20	1	ADB84282 Human mucolipin (M
4	16	4.0	16	1	ABE28963 Human MCOLN1 gene
C 5	15.4	3.8	17	1	ADB40824 Tumour suppression
6	14	3.5	15	1	ACD66351 Anti-HCV nucleic a
7	14	3.5	15	1	ACD66421 Anti-HCV enzymatic
8	14	3.5	15	1	ADI87738 Anti-HCV molecule

ALIGNMENTS

RESULT 1
ADU28709
ID ADU28709 standard; DNA; 21 BP.
XX AC
XX ADU28709;
XX AC
XX 27-JAN-2005 (first entry)
XX DE
XX Knock-down target sequence #2107.
XX ds; RNA production; protein production; drug development;
XX knock-down target.
XX Unidentified.
XX OS
XX WO2004094636-A1.
XX FN
XX 04-NOV-2004.
XX PD
XX 24-APR-2003; 2003WO-EP004362.

XX 24-APR-2003; 2003WO-EP004362.
XX (GALA-) GALAPAGOS GENOMICS NV.
XX (VSCH/) VAN DER SCHUEREN J.
XX Arts GJF, Lambrecht MJY, Djokic K, Clasen RJ, Mesic E;
XX Griffioen S, Bergs CJL;
XX WPI; 2004-775940/76.
XX New knockdown sequences, useful in lowering the amount of RNA and/or
XX protein production in cells used in drug development process.
XX Claim 11; SEQ ID NO 2123; 402pp; English.
XX The invention relates to a polynucleotide comprising an RNA sequence. The
XX polynucleotides, vector, libraries, and method are useful in lowering the
XX amount of RNA and/or protein production in cells used in drug development
XX process. The present sequence represents a knock-down target sequence.
XX Sequence 21 BP; 7 A; 6 C; 5 G; 3 T; 0 U; 0 Other;
XX
XX Query Match 5.2%; Score 21; DB 1; Length 21;
XX Best Local Similarity 100.0%; Pred. No. 0.71;
XX Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 6994 ACATCCAGGAGTGAAGCACC 7014
DB 1 ACATCCAGGAGTGAAGCACC 21
RESULT 2
AEB28960/c
ID AEB28960 standard; DNA; 21 BP.
AC AEB28960;
XX 22-SEP-2005 (first entry)
XX Human MCOLN1 gene PCR primer PRI R2 SEQ ID NO:4.
XX mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; PCR;
XX primer; ss.
XX Homo sapiens.
XX US2005153300-A1.
XX 14-JUL-2005. 10 17521110106
XX 09-JAN-2004; 2004US-00754446.
XX 09-JAN-2004; 2004US-00754446.
XX (QUES-) QUEST DIAGNOSTICS INC.
XX Sun W, Hantash F;
XX WPI; 2005-521160/53.
XX Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by
XX amplifying the nucleic acid, detecting amplified product with labeled
XX oligonucleotide probes via a change in fluorescence which indicates the
XX presence of an ML IV mutant.
XX Claim 1; SEQ ID NO 4; 15pp; English.
XX The invention relates to a method (M1) for detecting the presence of
XX mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1)
XX involves contacting the nucleic acid with oligonucleotide primers and
XX probes, conducting amplification by temperature cycling and monitoring
XX the accumulation of amplified nucleic acid by detecting an increase in

CC donor fluorophore fluorescence or decrease in acceptor fluorophore
 CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucolin-1
 CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a PCR primer for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

SQ Sequence 21 BP; 5 A; 7 C; 5 G; 4 T; 0 U; 0 Other;

Query Match 5.2%; Score 21; DB 1; Length 21;
 Best Local Similarity 100.0%; Pred. No. 0.71;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7017 AGTGTCTTCCAGCAGCGTGAG 7037

Db 21 AGTGTCTTCCAGCAGCGTGAG 1

RESULT 3

ID ADB84282/c
 ID ADB84282 standard; DNA; 20 BP.

XX AC ADB84282;

XX DT 04-DEC-2003 (first entry)

XX DE Human mucolin (MCOLIPIN-1) gene AJ haplotype analysis primer #4.

XX KW human; mucolin; MCOLN-1; mucolinipidosis IV; ion channel defect;

XX KW immunogen; gene therapy; ss; primer; PCR; AJ haplotype.

XX OS Homo sapiens.

XX PN US2003064363-A1.

XX PD 03-APR-2003.

XX PF 08-MAY-2001; 2001US-00851494.

XX PR 18-AUG-2000; 2000US-0226388P.

XX PA (MLFO-) ML4 FOUND & HAVARD COLLEGE.

XX PI Goldin E, Slaugenhaupt SA, Sun M, Acierno JS;

XX DR WPI; 2003-540779/51.

XX New mucolin (MCOLN1) polypeptides and nucleic acids, useful in gene
 PT therapy for treating mucolinipidosis IV or ion channel defect, or as
 PT immunogen to generate antibodies that recognize the MCOLN1 polypeptide.

PS Example; Page 18; 34pp; English.

XX The invention describes a nucleic acid (I) encoding a mucolin (MCOLN1)
 CC polypeptide, and sharing at least 95% sequence identity with sequence
 CC selected from 2 nucleotide sequences (designated S1 and S2, not given in
 CC the specification). A mutation of in the MCOLN1 gene encoding the MCOLN1
 CC polypeptide results in a defect in expression of a functional MCOLN1. The
 CC MCOLN1 gene is useful for the treatment of mucolinipidosis IV or
 CC ion channel defect. The MCOLN1 polypeptides are useful as immunogen to
 CC generate antibodies that recognize the MCOLN1 polypeptide. Such
 CC antibodies may be used for diagnostics and intracellular regulation of
 CC MCOLN1 activity. The polynucleotides may also be useful in gene therapy.
 CC This sequence represents a primer used to analyse the AJ haplotype of the
 CC human mucolin (MCOLN-1) gene.

XX SQ Sequence 20 BP; 5 A; 7 C; 5 G; 3 T; 0 U; 0 Other;

Query Match 5.0%; Score 20; DB 1; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.94;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7018 GTGTCTTCCAGCAGCGTGAG 7037

Db 20 GTGTCTTCCAGCAGCGTGAG 1

RESULT 4

AEB28963

ID AEB28963 standard; DNA; 16 BP.

XX AC AEB28963;

XX DT 22-SEP-2005 (first entry)

XX DE Human MCOLN1 gene probe DEL SEQ ID NO:7.

XX KW mucolinipidosis IV; high throughput screening; mucolinipin-1; MCOLN1; probe;
 XX ss.

XX OS Homo sapiens.

XX PN US2005153300-A1.

XX PD 14-JUL-2005.

XX PF 09-JAN-2004; 2004US-00754446.

XX PR 09-JAN-2004; 2004US-00754446.

XX PA (QUES-) QUEST DIAGNOSTICS INC.

XX PI Sun W, Hantash F;

XX DR WPI; 2005-521160/53.

XX Diagnosing mucolinipidosis (ML) IV mutant sequence in nucleic acids, by
 PT amplifying the nucleic acid, detecting amplified product with labeled
 PT oligonucleotide probes via a change in fluorescence which indicates the
 PT presence of an ML IV mutant.

PS Claim 3; SEQ ID NO 7; 15pp; English.

CC The invention relates to a method (M1) for detecting the presence of
 CC mucolinipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1)
 CC involves contacting the nucleic acid with oligonucleotide primers and
 CC probes, conducting amplification by temperature cycling and monitoring
 CC the accumulation of amplified nucleic acid by detecting an increase in
 CC donor fluorophore fluorescence or decrease in acceptor fluorophore
 CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucolin-1
 CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a probe for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

SQ Sequence 16 BP; 5 A; 7 C; 3 G; 1 T; 0 U; 0 Other;

Query Match 4.0%; Score 16; DB 1; Length 16;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6982 AGACCCAGGCCACAT 6997

Db 1 AGACCCAGGCCACAT 16

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 15:03:21 ; Search time 0.001 Seconds
(without alignments)
147.568 Million cell updates/sec

Title: af287270
Perfect score: 401
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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 0.5

Searched: 10 seqs, 184 residues

Total number of hits satisfying chosen parameters: 20

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 10 summaries

Database : af287270_copy_6956_7356.rnpbm4.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	25	6.2	25	1 US-11-060-756-186093	Sequence 186093,
2	25	6.2	25	1 US-11-060-756-261208	Sequence 261208,
C 3	21	5.2	21	1 US-10-754-446-4	Sequence 4, Appli
C 4	20	5.0	20	1 US-09-851-494B-11	Sequence 11, Appl
5	16	4.0	16	1 US-10-754-446-7	Sequence 7, Appli
C 6	14.4	3.6	16	1 US-09-896-324B-62	Sequence 62, Appl
C 7	14.4	3.6	16	1 US-10-236-363A-44	Sequence 44, Appl
8	14	3.5	15	1 US-09-740-332-4784	Sequence 4784, Ap
9	14	3.5	15	1 US-09-817-879-4784	Sequence 4784, Ap
10	14	3.5	15	1 US-10-669-841-7381	Sequence 7381, Ap

ALIGNMENTS

RESULT 1
US-11-060-756-186093
; Sequence 186093, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 186093
; LENGTH: 25
; TYPE: DNA
; ORGANISM: probe

US-11-060-756-186093

Query Match 6.2%; Score 25; DB 1; Length 25;
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Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6991 CCCACATCCAGGAGTGTAAAGCACCC 7015
Db 1 CCCACATCCAGGAGTGTAAAGCACCC 25

RESULT 2

US-11-060-756-261208
; Sequence 261208, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 261208
; LENGTH: 25
; TYPE: DNA
; ORGANISM: probe
US-11-060-756-261208

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Best Local Similarity 100.0%; Pred. No. 0.46;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6989 GGGCCACATCCAGGAGTGTAAAGCAC 7013
Db 1 GGGCCACATCCAGGAGTGTAAAGCAC 25

RESULT 3

US-10-754-446-4/c
; Sequence 4, Application US/10754446
; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: SUN, WEIMIN
; APPLICANT: HANTASH, FERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10/754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 3.2
; SEQ ID NO 4
; LENGTH: 21
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-10-754-446-4

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Best Local Similarity 100.0%; Pred. No. 1.2;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7017 AGTGTCTTCCAGCAGCGGTGAG 7037
Db 21 AGTGTCTTCCAGCAGCGGTGAG 1

RESULT 4

US-09-851-494B-11/c

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; Sequence 11, Application US/09851494B
; Publication No. US20030064363A1
; GENERAL INFORMATION:
; APPLICANT: ML4 Foundation
; APPLICANT: Goldin, Ehud
; APPLICANT: Slaugenhaupt, Susan A.
; APPLICANT: Sun, Mei
; APPLICANT: Aclerno, James S.
; TITLE OF INVENTION: A Gene Encoding A New TRP Channel is Mutated in Mucopolidosis IV
; FILE REFERENCE: 3394/1H557US1
; CURRENT APPLICATION NUMBER: US/09/851,494B
; CURRENT FILING DATE: 2002-07-12
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 11
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: PCR primer
US-09-851-494B-11

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Best Local Similarity 100.0%; Pred. No. 1.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7018 GTGCTTCCAGCGGTGAG 7037
Db 20 GTGCTTCCAGCGGTGAG 1

RESULT 5
US-10-754-446-7
; Sequence 7, Application US/10754446
; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: SUN, WEIMIN
; APPLICANT: HANTASH, FERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10/754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 3.2
; SEQ ID NO 7
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Probe
US-10-754-446-7

Query Match          4.0%; Score 16; DB 1; Length 16;
Best Local Similarity 100.0%; Pred. No. 4.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6982 AGACCCAGGCCACAT 6997
Db 1 AGACCCAGGCCACAT 16

RESULT 6
US-09-896-324B-62/c
; Sequence 62, Application US/09896324B
; Publication No. US20030148276A1
; GENERAL INFORMATION:
; APPLICANT: Li, Bi-yu
; TITLE OF INVENTION: METHOD FOR IDENTIFICATION, SEPARATION AND QUANTITATIVE MEASUREMENT
; FILE REFERENCE: 45163-1008
; CURRENT APPLICATION NUMBER: US/09/896,324B
; CURRENT FILING DATE: 2002-11-04
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; NUMBER OF SEQ ID NOS: 89
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 62
; LENGTH: 16
; TYPE: DNA
; ORGANISM: CD18-BsaJ I-ga
US-09-896-324B-62

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Best Local Similarity 93.8%; Pred. No. 5.9;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7176 GTCCTGTGCTGAGATC 7191
Db 16 GTCAGTGTCTGAGATC 1

RESULT 7
US-10-236-363A-44/c
; Sequence 44, Application US/10236363A
; Publication No. US20030165923A1
; GENERAL INFORMATION:
; APPLICANT: Li, Bi-yu
; APPLICANT: Wang, Xun
; APPLICANT: Shi, Liang
; TITLE OF INVENTION: METHOD FOR IDENTIFICATION OF GENETIC MARKERS
; FILE REFERENCE: TM0011-CIP
; CURRENT APPLICATION NUMBER: US/10/236,363A
; CURRENT FILING DATE: 2002-11-19
; PRIOR APPLICATION NUMBER: US 09/896,324
; PRIOR FILING DATE: 2001-06-29
; PRIOR APPLICATION NUMBER: US 60/215,596
; PRIOR FILING DATE: 2000-06-30
; NUMBER OF SEQ ID NOS: 139
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 44
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: adapter oligonucleotide
US-10-236-363A-44

Query Match          3.6%; Score 14.4; DB 1; Length 16;
Best Local Similarity 93.8%; Pred. No. 5.9;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 7176 GTCCTGTGCTGAGATC 7191
Db 16 GTCAGTGTCTGAGATC 1

RESULT 8
US-09-740-332-4784
; Sequence 4784, Application US/09740332
; Publication No. US20030125270A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals Inc.
; TITLE OF INVENTION: Enzymatic Nucleic Acid Treatment of Diseases or Conditions Related
; FILE REFERENCE: RPI 400/003
; CURRENT APPLICATION NUMBER: US/09/740,332
; CURRENT FILING DATE: 2001-03-26
; NUMBER OF SEQ ID NOS: 9704
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4784
; LENGTH: 15
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION:
; OTHER INFORMATION: oligonucleotide substrate
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 17, 2006, 15:04:33 ; Search time 0.001 Seconds
(without alignments)
63.358 Million cell updates/sec

Title: af287270
Perfect score: 401
Sequence: 1 CAGTGGCGGATCCCATCA.....CATCTTGGCCCTACCGCT 401

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 0.5

Searched: 4 seqs, 79 residues

Total number of hits satisfying chosen parameters: 8

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 4 summaries

Database : af287270_copy_6956_7356.rnpbn4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	20	5.0	21	1 US-10-310-914A-66074	Sequence 66074, A
2	18.4	4.6	20	1 US-10-310-914A-253105	Sequence 253105, A
c 3	18	4.5	20	1 US-10-310-914A-66019	Sequence 66019, A
4	17	4.2	18	1 US-10-310-914A-918753	Sequence 918753, A

ALIGNMENTS

RESULT 1
US-10-310-914A-66074/c
; Sequence 66074, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 66074
; LENGTH: 21
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-66074

Query Match 5.0%; Score 20; DB 1; Length 21;
Best Local Similarity 100.0%; Pred. No. 0.92; Indels 0; Gaps 0;
Matches 20; Conservative 0; Mismatches 0

Qy 7057 CACTGACCAGGGGCCCTGGC 7076
Db 20 CACTGACCAGGGGCCCTGGC 1

RESULT 2

US-10-310-914A-253105
; Sequence 253105, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 253105
; LENGTH: 20
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-253105

Query Match 4.6%; Score 18.4; DB 1; Length 20;
Best Local Similarity 90.0%; Pred. No. 1.5;
Matches 18; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 7038 CCCCTGAGCCCCAGACCAGC 7057
Db 1 CCCCTGAGCCCCAGCCCAGC 20

RESULT 3

US-10-310-914A-66019/c
; Sequence 66019, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 66019
; LENGTH: 20
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-66019

Query Match 4.5%; Score 18; DB 1; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7057 CACTGACCAGGGGCCCTG 7074
Db 18 CACTGACCAGGGGCCCTG 1

RESULT 4

US-10-310-914A-918753
; Sequence 918753, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01

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; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 918753
; LENGTH: 18
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-918753
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Query Match      4.2%; Score 17; DB 1; Length 18;
Best Local Similarity 94.1%; Pred. No. 2.4;
Matches 16; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
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Oy 7189 ATCCCCCAAGCCCCAGA 7205
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Db 2 AUCCCCCAAGCCCCAGA 18
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Title: af287270
Perfect score: 401
Sequence: 1 GGGTCTCAGCAGGACAA.....TAGTCTACTGAGCCTCGAC 401

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 0.5
Searched: 101 seqs, 1809 residues

Total number of hits satisfying chosen parameters: 202

Minimum DB seq length: 15
Maximum DB seq length: 30

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 101 summaries

Database : af287270_copy_100_500.rge4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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2	20.4	5.1	22	1	AR076805
3	20.4	5.1	22	1	AR242947
4	20.4	5.1	22	1	AR345130
5	20.4	5.1	22	1	AX384999
6	20.0	5.0	20	1	BD233827
7	20.0	5.0	20	1	AR321577
8	20.0	5.0	20	1	AR542568
9	19.4	4.8	21	1	AR345126
10	19.4	4.8	21	1	AX741051
11	19.0	4.7	19	1	CS010199
12	19.0	4.7	19	1	CS028041
13	19.0	4.7	19	1	CS028508
14	19.0	4.7	19	1	AX114983
15	19.0	4.7	20	1	AR152875
16	19.0	4.7	20	1	BD134331
17	19.0	4.7	20	1	BD138317
18	19.0	4.7	20	1	ARG30295
19	19.0	4.7	20	1	ARG34606
20	18.4	4.6	20	1	AR162414
21	18.4	4.6	20	1	BD138315
22	18.4	4.6	20	1	BD138316
23	18.4	4.6	20	1	I31362
24	18.4	4.6	20	1	AR323228
25	18.4	4.6	20	1	AR370243
26	18.2	4.5	19	1	AR074597
27	18.2	4.5	19	1	AR083936
28	18.2	4.5	19	1	I23816
29	18.2	4.5	19	1	I29970
30	18.0	4.5	18	1	AR576394
31	18.0	4.5	19	1	CS092555
32	18.0	4.5	19	1	CS092721
33	18.0	4.5	19	1	AX133851

ALIGNMENTS

ACCESSION:AR148945	1	19	4.3	17.4	34
ACCESSION:CS027943	1	19	4.3	17.4	35
ACCESSION:CS028410	1	19	4.3	17.4	c 36
ACCESSION:AX116706	1	19	4.3	17.4	c 37
ACCESSION:AR044034	1	17	4.2	17.0	c 38
ACCESSION:AR592720	1	17	4.2	17.0	c 39
ACCESSION:AX571818	1	17	4.2	17.0	c 40
ACCESSION:AX761572	1	17	4.2	17.0	41
ACCESSION:AR576395	1	18	4.2	17.0	42
ACCESSION:AR576396	1	18	4.2	17.0	43
ACCESSION:AR585363	1	18	4.2	17.0	44
ACCESSION:AR612296	1	18	4.2	17.0	45
ACCESSION:AR612299	1	18	4.2	17.0	c 46
ACCESSION:AR637194	1	18	4.2	17.0	47
ACCESSION:AR154019	1	18	4.1	16.4	c 48
ACCESSION:BD130125	1	18	4.1	16.4	c 49
ACCESSION:AR565243	1	18	4.1	16.4	c 50
ACCESSION:AR575575	1	18	4.1	16.4	51
ACCESSION:AB069644	1	18	4.1	16.4	52
ACCESSION:AR633675	1	18	4.0	16.2	53
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ACCESSION:AR614733	1	17	4.0	16.0	59
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ACCESSION:AX692568	1	17	4.0	16.0	63
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ACCESSION:BD202918	1	17	3.8	15.4	67
ACCESSION:AR597131	1	17	3.8	15.4	c 68
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ACCESSION:AX729877	1	17	3.8	15.4	c 74
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ACCESSION:AX734071	1	17	3.8	15.4	c 79
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ACCESSION:AX736964	1	17	3.8	15.4	c 81
ACCESSION:AX737636	1	17	3.8	15.4	c 82
ACCESSION:AX737828	1	17	3.8	15.4	c 83
ACCESSION:AX738556	1	17	3.8	15.4	c 84
ACCESSION:AX739093	1	17	3.8	15.4	c 85
ACCESSION:AX758145	1	17	3.8	15.4	c 86
ACCESSION:AX760525	1	17	3.8	15.4	c 87
ACCESSION:AX761540	1	17	3.8	15.4	c 88
ACCESSION:CQ971639	1	15	3.7	15.0	89
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ACCESSION:CS002308	1	15	3.7	15.0	91
ACCESSION:CS002310	1	15	3.7	15.0	92
ACCESSION:CS048833	1	15	3.7	15.0	c 93
ACCESSION:CS074138	1	15	3.7	15.0	c 94
ACCESSION:AR561375	1	15	3.7	15.0	95
ACCESSION:AR612295	1	15	3.7	15.0	96
ACCESSION:AR612298	1	15	3.7	15.0	c 97
ACCESSION:AR630722	1	15	3.7	15.0	98
ACCESSION:AR436001	1	16	3.6	14.4	99
ACCESSION:CS002304	1	15	3.5	14.0	100
ACCESSION:CS002312	1	15	3.5	14.0	101

RESULT 1
LOCUS AX614112 25 bp DNA linear PAT 17-FEB-2003
DEFINITION Sequence 5137 from Patent WO02072882.
ACCESSION AX614112
VERSION AX614112.1 GI:28409541
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1
AUTHORS Cullen, P. and Seedorf, U.
TITLE Coronary chip
JOURNAL Patent: WO 02072882-A 5137 19-SEP-2002;
OGHAM GmbH (DE)
FEATURES
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
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Best Local Similarity 96.0%; Pred. No. 8.9;
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Qy 444 GCTCTGTGCCCCAGCGTGGAGTGCA 468
Db 1 GCTCTGTGCCCCAGCGTGGAGTGCA 25
RESULT 2
LOCUS AR076805 22 bp DNA linear PAT 30-AUG-2000
DEFINITION Sequence 2 from patent US 5959171.
ACCESSION AR076805
VERSION AR076805.1 GI:10003551
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 22)
AUTHORS Hyttinen, J.-M., Korhonen, V.-P., and Janne, J.
TITLE Method for the production of biologically active polypeptides in a mammal's
JOURNAL Patent: US 5959171-A 2 28-SEP-1999;
FEATURES
source Location/Qualifiers
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/organism="unknown"
/mol_type="unassigned DNA"
Query Match 5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 16;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 464 GTGCAGTGGTGTCATCAGCT 485
Db 1 GTGCAGTGGTGTCATCAGCT 22
RESULT 3
LOCUS AR242947 22 bp DNA linear PAT 20-DEC-2002
DEFINITION Sequence 93 from patent US 6475739.
ACCESSION AR242947
VERSION AR242947.1 GI:27289609
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 22)
AUTHORS Brunkow, M.E., Proll, S., Paepel, B. and Staehling-Hampton, K.

TITLE Methods for identifying genomic deletions
JOURNAL Patent: US 6475739-A 93 05-NOV-2002;
Celltech R&D, Inc.; Bothell, WA
FEATURES
source Location/Qualifiers
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/organism="unknown"
/mol_type="genomic DNA"
Query Match 5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 16;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 455 CAGCGTGGAGTGGAGTGGTGTG 476
Db 1 CAGCGTGGAGTGGAGTGGTGTG 22
RESULT 4
LOCUS AR345130 22 bp DNA linear PAT 17-AUG-2003
DEFINITION Sequence 11 from patent US 6583112.
ACCESSION AR345130
VERSION AR345130.1 GI:33741766
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 22)
AUTHORS Fu, Y.-H., Yu, C.-E., Oshima, J., Mulligan, J.T. and Schellenberg, G.D.
TITLE Gene products related to Werner's syndrome
JOURNAL Patent: US 6583112-A 11 24-JUN-2003;
FEATURES
source Location/Qualifiers
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/organism="unknown"
/mol_type="genomic DNA"
Query Match 5.1%; Score 20.4; DB 1; Length 22;
Best Local Similarity 95.5%; Pred. No. 16;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 463 AGTGCAGTGGTGTCATCAGC 484
Db 1 AGTGCAGTGGTGTCATCAGC 22
RESULT 5
LOCUS AX384999 22 bp DNA linear PAT 19-MAR-2002
DEFINITION Sequence 93 from Patent WO0210455.
ACCESSION AX384999
VERSION AX384999.1 GI:19578127
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Brunkow, M.E., Proll, S. and Paepel, B.
TITLE Methods for identifying genomic deletions
JOURNAL Patent: WO 0210455-A 93 07-FEB-2002;
Celltech R & D, Inc. (US); Straehling-Hampton, Karen (US)
FEATURES
source Location/Qualifiers
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/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="PCR primer"
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Best Local Similarity 95.5%; Pred. No. 16;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 455 CAGCGTGGAGTGGAGTGGTGTG 476

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 736.68 Seconds
(without alignments)
1157.425 Million cell updates/sec

Title: US-10-754-446-6

Perfect score: 15

Sequence: 1 ctgcccacggtacct 15

Scoring table:

IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 5883141 seqs, 28421725653 residues

Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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1: gb_ba.*

2: gb_in.*

3: gb_env.*

4: gb_on.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

8: gb_pr.*

9: gb_ro.*

10: gb_sts.*

11: gb_sy.*

12: gb_un.*

13: gb_vi.*

14: gb_htg.*

15: gb_pl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	15	100.0	194763	14	AC150852
5	15	100.0	216844	14	AC130788
6	15	100.0	218893	14	AC150547
7	15	100.0	221544	14	AC127862
8	15	100.0	224638	14	AC135282
9	15	100.0	235008	14	AC103239
10	15	100.0	239553	14	AC123201
11	15	100.0	246860	14	AC115225
12	15	100.0	320746	14	AC128373
13	14	93.3	424	10	BV105141
14	14	93.3	807	6	BD162787
15	14	93.3	807	6	AX120670
16	14	93.3	930	6	AX064573
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19	14	93.3	1486	15	YSPADH	J01341	fishon yeas
20	14	93.3	1985	15	AB000261	AB000261	Aspergill
21	14	93.3	3369	15	AB000262	AB000262	Aspergill
22	14	93.3	7319	6	AR227050	AR227050	Sequence
23	14	93.3	9999	15	SPCC13B11	AL032681	S.pombe c
24	14	93.3	11304	1	AE012358	AE012358	Xanthomon
25	14	93.3	12403	1	AE001706	AE001706	Thermotog
26	14	93.3	62398	14	AC166844	AC166844	Bos tauru
27	14	93.3	69517	14	AC036117	AC036117	Homo sapi
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32	14	93.3	110000	1	BA000036_05	Continuation (6 of	
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36	14	93.3	110000	15	AE017342_11	Continuation (12 o	
37	14	93.3	110000	15	AE017342_12	Continuation (13 o	
38	14	93.3	110000	15	AE017348_02	Continuation (3 of	
39	14	93.3	116409	9	AC131118	AC131118	Mus muscu
40	14	93.3	139697	14	AC134207	AC134207	Rattus no
41	14	93.3	140176	8	AC022188	AC022188	Homo sapi
42	14	93.3	140219	9	AL603714	AL603714	Mouse DNA
43	14	93.3	143509	14	AC068683	AC068683	Homo sapi
44	14	93.3	143733	14	AC152465	AC152465	Dasybus n
45	14	93.3	155628	14	AC027810	AC027810	Homo sapi

ALIGNMENTS

RESULT 1
LOCUS CQ750386
DEFINITION Sequence 36320 from Patent WO02068579.
ACCESSION CQ750386
VERSION CQ750386.1 GI:42381467
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE 1
AUTHORS Venter, C.J., Adams, M.C., Li, P.W. and Myers, E.W.
TITLE Kits, such as nucleic acid arrays, comprising a majority of humanexons or transcripts, for detecting expression and other uses thereof
JOURNAL Patent: WO 02068579-A 36320 06-SEP-2002;
PE Corporation (NY) (US)
FEATURES
source Location/Qualifiers
1. 1935
/organism="Homo sapiens"
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/db_xref="taxon:9606"

Query Match 100.0%; Score 15; DB 6; Length 1935;
Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTGCCCCACGGTACCT 15
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Db 912 CTGCCCCACGGTACCT 926

RESULT 2
LOCUS AC130385
DEFINITION Homo sapiens chromosome 17 clone RP13-49B7 map 17, LOW-PASS
SEQUENCE SAMPLING.
ACCESSION AC130385

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VERSION AC130385.4 GI:23683263
KEYWORDS HTG; HTGS PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1 (bases 1 to 65347)
AUTHORS Birren,B., Nusbaum,C. and Lander,E.
TITLE Homo sapiens chromosome 17, clone RP13-49B7
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 65347)
AUTHORS Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulne,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., MacLean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schuback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (09-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
JOURNAL 3 (bases 1 to 65347)
REFERENCE 3 (bases 1 to 65347)
AUTHORS Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulne,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., MacLean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schuback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (10-OCT-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
JOURNAL On Oct 10, 2002 this sequence version replaced gi:23322704.
COMMENT All repeats were identified using RepeatMasker:
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L27996
Center clone name: 49_B_7
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* NOTE: This record contains 83 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for

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* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
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* 681: contig of 681 bp in length
* 682
* 781: gap of 100 bp
* 782
* 1477: contig of 696 bp in length
* 1478
* 1577: gap of 100 bp
* 1578
* 2269: contig of 692 bp in length
* 2270
* 2369: gap of 100 bp
* 2370
* 3056: contig of 687 bp in length
* 3057
* 3156: gap of 100 bp
* 3157
* 3835: contig of 679 bp in length
* 3836
* 3935: gap of 100 bp
* 3936
* 4629: contig of 694 bp in length
* 4630
* 4729: gap of 100 bp
* 4730
* 5426: contig of 697 bp in length
* 5427
* 5526: gap of 100 bp
* 5527
* 6218: contig of 692 bp in length
* 6219
* 6318: gap of 100 bp
* 6319
* 7003: contig of 685 bp in length
* 7004
* 7103: gap of 100 bp
* 7104
* 7798: contig of 695 bp in length
* 7799
* 7898: gap of 100 bp
* 7899
* 8584: contig of 686 bp in length
* 8585
* 8684: gap of 100 bp
* 8685
* 9370: contig of 686 bp in length
* 9371
* 9470: gap of 100 bp
* 9471
* 10159: contig of 689 bp in length
* 10160
* 10259: gap of 100 bp
* 10260
* 10954: contig of 695 bp in length
* 10955
* 11054: gap of 100 bp
* 11055
* 11740: contig of 686 bp in length
* 11741
* 11840: gap of 100 bp
* 11841
* 12487: contig of 647 bp in length
* 12488
* 12587: gap of 100 bp
* 12589
* 13282: contig of 695 bp in length
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* 13382: gap of 100 bp
* 13383
* 14089: contig of 687 bp in length
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* 14169: gap of 100 bp
* 14170
* 14855: contig of 686 bp in length
* 14856
* 14955: gap of 100 bp
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* 15633: contig of 678 bp in length
* 15634
* 15733: gap of 100 bp
* 15734
* 16426: contig of 693 bp in length
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* 16526: gap of 100 bp
* 16527
* 17198: contig of 672 bp in length
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* 17298: gap of 100 bp
* 17299
* 17983: contig of 685 bp in length
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* 18777: contig of 694 bp in length
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* 18877: gap of 100 bp
* 18878
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* 19648: gap of 100 bp
* 19649
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* 20445: gap of 100 bp
* 20446
* 21138: contig of 693 bp in length
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* 21238: gap of 100 bp
* 21239
* 21942: contig of 704 bp in length
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* 22042: gap of 100 bp
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* 22751: contig of 709 bp in length
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* 23545: contig of 694 bp in length
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* 23645: gap of 100 bp
* 23646
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* 25092: contig of 662 bp in length
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* 25893: contig of 701 bp in length
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* 25993: gap of 100 bp
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* 26679: contig of 686 bp in length

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Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 1677.48 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-6

Perfect score: 15

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Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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2: gb_est2:*
3: gb_est3:*
4: gb_hc:*
5: gb_est4:*
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7: gb_est6:*
8: gb_est7:*
9: gb_gses1:*
10: gb_gses2:*
11: gb_gses3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	15	100.0	398	6	CB707082 AMGNNUC:N
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5	15	100.0	680	7	CK845557 UI-R-BJ1
6	15	100.0	696	8	DN105468 1101440 M
7	15	100.0	697	8	CV796826 UI-R-EB1
8	15	100.0	748	7	CO565750 AGENCOURT
9	15	100.0	763	8	CV991014 IPCGPr1.5
10	15	100.0	905	7	CN248276 EST014180
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12	14	93.3	303	2	BF394984 UI-R-CM0
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16	14	93.3	373	5	BU994782 HM08C24r
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18	14	93.3	445	1	AU007564 AU007564
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22	14	93.3	475	1	AW342926 fj80910.y

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	25	14	93.3	492	3	BI538165	BI538165
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	27	14	93.3	515	6	CF364476	CF364476
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	32	14	93.3	547	3	BI343534	BI343534
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ALIGNMENTS

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DEFINITION BF353711
ACCESSION BF353711
VERSION BF353711.1 GI:11312785
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 332)
AUTHORS Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R., Ngai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldman, G.H., Carvalho, A.F., Matekuma, A., Baia, G.S., Simpson, D.H., Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.
TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
PUBMED 10737800
COMMENT Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Frudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=QV2&t2=QV2-HT0698-110700-267-a01&t3=2000-07-11&t4=1)
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High quality sequence stop: 331.
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ORIGIN

Query Match 100.0%; Score 15; DB 2; Length 332;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTGCCACCGTACTT 15
Db 236 CTGCCACCGTACTT 250

RESULT 2

CB707082 398 bp mRNA linear EST 10-APR-2003
LOCUS AMGNNUC.NRHV5-00217-H7-A W Rat hypothalamus (10471) Rattus
DEFINITION norvegicus cDNA clone nrhys-00217-h7 5', mRNA sequence.

ACCESSION CB707082
VERSION CB707082.1 GI:29764230
KEYWORDS EST.

SOURCE

Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridea; Muridae; Murinae; Rattus.

1 (bases 1 to 398)

Angen EST Program.

Angen Rat EST Program

Unpublished (2003)

Contact: Dan Fitzpatrick

Angen, Inc

One Angen Center Drive, Thousand Oaks, CA 91320-1799, USA

Tel: 805 447-4881

Plate: 00217 row: h column: 7.

FEATURES

source

1. .398

/organism="Rattus norvegicus"

/mol_type="mRNA"

/db_xref="taxon:10116"

/clone="nrhys-00217-h7"

/clone_lib="W Rat hypothalamus (10471)"

/notes="Vector: pSPORT1; Site 1: Sall; Site 2: NotI; W Rat hypothalamus adult female Wistar rat avg. insert size 2.3 kb fraction 6 and 7"

ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 398;
Best Local Similarity 100.0%; Pred. No. 1.7e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTGCCACCGTACTT 15
Db 40 CTGCCACCGTACTT 54

RESULT 3

CB579676 615 bp mRNA linear EST 03-APR-2003
LOCUS AMGNNUC.NRHV5-00106-F3-A W Rat hypothalamus (10471) Rattus
DEFINITION norvegicus cDNA clone nrhys-00106-f3 5', mRNA sequence.

ACCESSION CB579676
VERSION CB579676.1 GI:29523717
KEYWORDS EST.

SOURCE

Rattus norvegicus (Norway rat)

ORGANISM

Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;

Sciurognathi; Muridea; Muridae; Murinae; Rattus.
1 (bases 1 to 615)

Angen EST Program.

Angen Rat EST Program

Unpublished (2003)

Contact: Dan Fitzpatrick

Angen, Inc

One Angen Center Drive, Thousand Oaks, CA 91320-1799, USA

Tel: 805 447-4881

Plate: 00106 row: f column: 3.

FEATURES

source

1. .615

/organism="Rattus norvegicus"

/mol_type="mRNA"

/db_xref="taxon:10116"

/clone="nrhys-00106-f3"

/clone_lib="W Rat hypothalamus (10471)"

/note="Vector: pSPORT1; Site 1: Sall; Site 2: NotI; W Rat hypothalamus adult female Wistar rat avg. insert size 2.3 kb fraction 6 and 7"

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTGCCACCGTACTT 15
Db 145 CTGCCACCGTACTT 159

RESULT 4

CG897726 674 bp DNA linear GSS 08-DEC-2003
LOCUS pastbac058xx02.bl.ab1 Res147 1 Pasteuria penetrans genomic 5',
DEFINITION genomic survey sequence.

ACCESSION CG897726

VERSION CG897726.1 GI:39553221

KEYWORDS GSS.

SOURCE Pasteuria penetrans

ORGANISM

Pasteuria penetrans

Bacteria; Firmicutes; Bacillales; Alicyclobacillaceae; Pasteuria.

1 (bases 1 to 674)

Opperman,C.H., Davies,K.G., Sosinski,B.R., Waterman,J. and Burke,M.

Unpublished Data

Unpublished (2003)

Contact: Opperman CH

Center for the Biology of Nematode Parasitism and Nematode

Interactions Unit

North Carolina State University and Rothamsted Research, Ltd.

Box 7253, NCSU, Raleigh, NC 27606, USA

Email: warthog@unity.ncsu.edu

Homology: e-val = 1e-51. Description = DNA ligase

(polydeoxyribonucleotide synthase [NAD+])

gi|3688229|emb|CAA09732.1| DNA ligase [Geobacillus

stearothermophilus] Homology: e-val = 1e-50. Description = DNA

ligase (polydeoxyribonucleotide synthase (NAD+)) [Bacillus

halodurans] gi|15213980|sp|Q9KF37|DNLJ_BACHD DNA ligase

(polydeoxyribonucleotide synthase [NAD+]) gi|25293998|pir|A83731

DNA ligase (polydeoxyribonucleotide synthase (NAD+)) lig [imported]

- row: k column: 2

Class: shotgun

High quality sequence stop: 674.

FEATURES

source

1. .674

/organism="Pasteuria penetrans"

/mol_type="genomic DNA"

/strain="Res147"

/db_xref="taxon:86005"

/clone_lib="Res147 1"

ORIGIN

Query Match 100.0%; Score 15; DB 10; Length 674;

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 203.52 Seconds
(without alignments)
491.207 Million cell updates/sec

Title: US-10-754-446-6

Perfect score: 15

Sequence: 1 ctgccacggtacct 15

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

N_Geneseq_21.*
1: geneseqn1980s.*
2: geneseqn1990s.*
3: geneseqn2000s.*
4: geneseqn2001as.*
5: geneseqn2001bs.*
6: geneseqn2002as.*
7: geneseqn2002bs.*
8: geneseqn2003as.*
9: geneseqn2003bs.*
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11: geneseqn2003ds.*
12: geneseqn2004as.*
13: geneseqn2004bs.*
14: geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	15	100.0	15	14 AEB28962	Aeb28962 Human MCO
2	14	93.3	560	13 ADQ49258	Adq49258 Novel can
C 3	14	93.3	807	5 AAH65551	Aah65551 C glutami
C 4	14	93.3	930	4 AAF72180	Aaf72180 Coryneb
C 5	14	93.3	937	10 ADD13464	Add13464 C. glutam
6	14	93.3	1050	13 ADT47984	Adt47984 Bacterial
7	14	93.3	1365	10 ADC81562	Adc81562 Recombina
8	14	93.3	1422	13 ADS48605	Ads48605 Bacterial
9	14	93.3	1524	13 ADS56578	Ads56578 Bacterial
10	14	93.3	34980	5 AAH68525	Aah68525 C glutami
11	13.4	89.3	16	14 AEB28961	Aeb28961 Human MCO
C 12	13.4	89.3	118	5 ABV43024	Abv43024 Human pro
C 13	13.4	89.3	118	5 ABV34163	Abv34163 Human pro
C 14	13.4	89.3	146	12 ACH93483	Ach93483 Human gen
C 15	13.4	89.3	252	12 ADP61947	Adp61947 Maize car
C 16	13.4	89.3	265	12 ADP61965	Adp61965 Maize car
C 17	13.4	89.3	267	12 ADP61934	Adp61934 Maize car
C 18	13.4	89.3	270	12 ADP61924	Adp61924 Maize car
C 19	13.4	89.3	272	12 ADP61945	Adp61945 Maize car

C 20	13.4	89.3	273	2 AAV09248	Aav09248 Human cyt
C 21	13.4	89.3	273	2 AAV12206	Aav12206 Human ret
C 22	13.4	89.3	273	6 AAD24507	Aad24507 Human P45
C 23	13.4	89.3	273	13 ADU66854	Adu66854 hp450RAI
C 24	13.4	89.3	273	14 ADV90794	Adv90794 Human ret
C 25	13.4	89.3	273	14 ADV62611	Adv62611 Human P45
C 26	13.4	89.3	274	2 AAV09249	Aav09249 Murine cy
C 27	13.4	89.3	274	2 AAV12207	Aav12207 Mouse ret
C 28	13.4	89.3	274	6 AAD24508	Aad24508 Mouse P45
C 29	13.4	89.3	274	13 ADU66855	Adu66855 mp450RAI
C 30	13.4	89.3	274	14 ADV90795	Adv90795 Murine re
C 31	13.4	89.3	274	14 ADV62612	Adv62612 Mouse P45
C 32	13.4	89.3	278	12 ADP61931	Adp61931 Maize car
C 33	13.4	89.3	285	12 ADP61916	Adp61916 Maize car
C 34	13.4	89.3	296	12 ADP61894	Adp61894 Maize car
C 35	13.4	89.3	298	4 AAL11859	Aal11859 Human bre
C 36	13.4	89.3	304	12 ADP61910	Adp61910 Maize car
C 37	13.4	89.3	311	12 ADP61878	Adp61878 Maize car
C 38	13.4	89.3	312	12 ADP61883	Adp61883 Maize car
C 39	13.4	89.3	314	12 ADP61875	Adp61875 Maize car
C 40	13.4	89.3	315	12 ADP61873	Adp61873 Maize car
C 41	13.4	89.3	321	12 ADP61869	Adp61869 Maize car
C 42	13.4	89.3	324	12 ADP61867	Adp61867 Maize car
C 43	13.4	89.3	341	4 AAS35653	Aas35653 Human car
C 44	13.4	89.3	341	10 ADE45732	Ade45732 Human car
C 45	13.4	89.3	341	13 ADJ07150	Adj07150 Human car

ALIGNMENTS

RESULT 1

AEB28962

ID AEB28962 standard; DNA; 15 BP.

XX AC AEB28962;

XX DT 22-SEP-2005 (first entry)

XX DE Human MCOLN1 gene probe IVS MUT SEQ ID NO:6.

XX KW mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; probe;

XX OS Homo sapiens.

XX PN US2005153300-A1.

XX PD 14-JUL-2005.

XX PF 09-JAN-2004; 2004US-00754446.

XX PR 09-JAN-2004; 2004US-00754446.

XX PA (QUES-) QUEST DIAGNOSTICS INC.

XX PI Sun W, Kantash F;

XX DR WPI; 2005-521160/53.

XX PT Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by

amplifying the nucleic acid, detecting amplified product with labeled

oligonucleotide probes via a change in fluorescence which indicates the

presence of an ML IV mutant.

XX PS Claim 3; SEQ ID NO 6; 15pp; English.

XX CC The invention relates to a method (M1) for detecting the presence of

mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1)

involves contacting the nucleic acid with oligonucleotide primers and

probes, conducting amplification by temperature cycling and monitoring

the accumulation of amplified nucleic acid by detecting an increase in

donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucopolin-1
 CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a probe for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

XX
 SQ Sequence 15 BP; 2 A; 7 C; 3 G; 3 T; 0 U; 0 Other;
 Query Match 100.0%; Score 15; DB 14; Length 15;
 Best Local Similarity 100.0%; Pred. No. 3.4e+02; Mismatches 0; Indels 0; Gaps 0;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTGCCACGGTACCT 15
 Db 1 CTGCCACGGTACCT 15
 |||||
 |||||

RESULT 2
 ADQ49258
 ID ADQ49258 standard; DNA; 560 BP.
 AC ADQ49258;
 XX
 XX 21-OCT-2004 (first entry)
 XX Novel canine microarray-related DNA sequence SeqID560.
 XX canine microarray; drug screening; toxicity assay;
 XX environmental pollutant; cellular response; gene expression profile;
 KW toxic response; liver necrosis; fatty liver disease;
 KW protein adduct formation; hepatitis; dog; ds.
 XX
 XX Canis familiaris.
 XX
 XX WO2004063324-A2.
 PN
 PD 29-JUL-2004.
 XX
 XX 05-MAY-2003; 2003WO-US013853.
 XX
 XX 03-MAY-2002; 2002US-0377240P.
 PR
 XX (GENE-) GENE LOGIC INC.
 PA (PFIZ) PFIZER PROD INC.
 XX
 XX Diggins JC, Porter M, Wei T;
 PI
 XX WPI; 2004-561890/54.
 DR
 XX New isolated nucleic acid molecule, useful for drug screening and
 XX toxicity assays or for assessing the impact, including toxicity, of a
 PT compound, pharmaceutical agent or environmental pollutant on a cell or
 PT living organism.
 XX
 XX Claim 1; SEQ ID NO 560; 41pp; English.
 PS
 XX This invention is related to a novel isolated canine nucleic acid
 CC sequences and the construction of canine microarrays containing a
 CC significant portion of the canine genome. The isolated canine nucleic
 CC acid sequences of the invention may be useful for drug screening and
 CC toxicity assays. The invention is therefore useful for assessing the
 CC impact, including toxicity, of a compound, pharmaceutical agent or
 CC environmental pollutant on a cell or living organism. The methods are
 CC useful for detecting genes that are up- or down-regulated in canines in a
 CC disease state. The sequences are useful as diagnostic agents or markers
 CC to detect a cellular response in a sample individually or as part of a
 CC gene expression profile. It is also useful as a target for agents that

CC modulate gene expression or activity. The database is useful for
 CC producing electronic Northern blots that allow the user to determine the cell
 CC type or tissue in which a given gene is expressed and to allow
 CC determination of the abundance or expression level of a given gene in a
 CC particular tissue or cell. The methods are useful for determining the
 CC similarity of a toxic response to one or more individual compounds. The
 CC methods are useful for predicting at least one toxic response or the
 CC likelihood that a compound or test agent will induce various specific
 CC pathologies such as those of the liver (liver necrosis, fatty liver
 CC disease, protein adduct formation or hepatitis), those of the kidney,
 CC heart, brain or testes, or other pathologies associated with at least one
 CC of the toxins. The methods are also useful for predicting or elucidating
 CC the potential cellular pathways influenced, induced or modulated by the
 CC compound or test agent due to the similarity of the expression profile
 CC compared to the profile induced by a known toxin. The present sequence is
 CC that of a canine DNA sequence which was claimed for use during the
 CC production of a canine microarray of the invention.

XX
 SQ Sequence 560 BP; 133 A; 168 C; 152 G; 102 T; 0 U; 5 Other;
 Query Match 93.3%; Score 14; DB 13; Length 560;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03; Mismatches 0; Indels 0; Gaps 0;
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTGCCACGGTACC 14
 Db 61 CTGCCACGGTACC 74
 |||||
 |||||

RESULT 3
 AAH65551/c
 ID AAH65551 standard; DNA; 807 BP.
 XX
 AC AAH65551;
 XX
 XX 26-SEP-2001 (first entry)
 DT
 XX C glutamicum coding sequence fragment SEQ ID NO: 586.
 DE
 XX Corynebacterium; amino acid synthesis; vitamin; saccharide;
 KW organic acid synthesis; ds.
 XX
 XX Corynebacterium glutamicum.
 OS
 XX EP1108790-A2.
 PN
 XX 20-JUN-2001.
 PD
 XX 18-DEC-2000; 2000EP-00127688.
 XX
 XX 16-DEC-1999; 99JP-00377484.
 PR
 XX 07-APR-2000; 2000JP-00159162.
 PR
 XX 03-AUG-2000; 2000JP-00280988.
 PR
 XX (KYOW) KYOWA HAKKO KOGYO KK.
 PA
 XX Nakagawa S, Mizoguchi H, Ando S, Havaashi M, Ochiai K, Yokoi H;
 PI Tateishi N, Senoh A, Ikeda M, Ozaki A;
 PI
 XX WPI; 2001-376931/40.
 DR
 XX P-PSDB; AAG90332.
 DR
 XX Novel polynucleotides derived from Corynebacterium, for identifying
 PT mutation point of a gene, measuring expression of a gene, analyzing
 PT expression profile or pattern of a gene and identifying homologous gene.
 XX
 XX Claim 8; SEQ ID NO 586; 246pp + Sequence Listing; English.
 XX
 XX The present invention provides a number of nucleotide and protein
 CC sequences from the Corynebacterium glutamicum. These
 CC are useful for identifying the mutation point of a gene derived from a
 CC mutant of corynebacterium, measuring expression amount and analysing
 CC the expression profile or expression pattern of a gene derived from

Result No.	Score	Query Match	Length	DB	ID	Description	
1	15	100.0	15	9	US-10-754-446-6	Sequence 6, Appli	
2	14	93.3	25	8	US-10-719-900-215409	Sequence 215409, A	
3	14	93.3	491	7	US-10-423-599-52366	Sequence 52366, A	
4	14	93.3	500	4	US-09-925-065A-757224	Sequence 757224, A	
5	14	93.3	807	3	US-09-738-626-586	Sequence 586, App	
6	14	93.3	937	8	US-10-494-675-31	Sequence 31, Appl	
7	14	93.3	1050	6	US-10-369-493-46422	Sequence 46422, A	
8	14	93.3	1365	6	US-10-372-473-8	Sequence 8, Appli	
9	14	93.3	1422	6	US-10-369-493-27035	Sequence 27035, A	
10	14	93.3	1524	6	US-10-369-493-32252	Sequence 32252, A	
11	14	93.3	7319	5	US-10-194-163-510	Sequence 510, App	
12	14	93.3	3309400	3	US-09-738-626-1	Sequence 1, Appli	
13	13.4	89.3	16	9	US-10-754-446-5	Sequence 5, Appli	
14	13.4	89.3	25	10	US-11-036-317-515390	Sequence 515390, A	
15	13.4	89.3	118	8	US-10-357-930-34181	Sequence 34181, A	
16	13.4	89.3	118	8	US-10-357-930-43043	Sequence 43043, A	
17	13.4	89.3	146	6	US-10-029-386-26678	Sequence 26678, A	
18	13.4	89.3	201	8	US-10-719-993-9298	Sequence 9298, Ap	
19	13.4	89.3	201	8	US-10-741-600-14149	Sequence 14149, A	
20	13.4	89.3	201	8	US-10-741-600-14194	Sequence 14194, A	
21	13.4	89.3	201	8	US-10-741-600-14264	Sequence 14264, A	
22	13.4	89.3	201	8	US-10-741-600-14300	Sequence 14300, A	
23	13.4	89.3	201	8	US-10-741-600-14338	Sequence 14338, A	

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; TYPE: DNA
; ORGANISM: Mus musculus
US-10-719-900-215409

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Best Local Similarity 100.0%; Pred. No. 9e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  2 TGCCACGGTACCT 15
    |||||
Db   3 TGCCACGGTACCT 16

RESULT 3
US-10-424-599-52366/c
; Sequence 52366, Application US/10424599
; Publication No. US20040031072A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa Thomas J
; APPLICANT: Kovalic David K
; APPLICANT: Zhou Yihua
; APPLICANT: Cao Yongwei
; TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated With
; FILE REFERENCE: 38-21(53223)B
; CURRENT APPLICATION NUMBER: US/10/424,599
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 285684
; SEQ ID NO 52366
; LENGTH: 491
; TYPE: DNA
; ORGANISM: Glycine max
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MRT3847_182C.1
US-10-424-599-52366

Query Match      93.3%; Score 14; DB 7; Length 491;
Best Local Similarity 100.0%; Pred. No. 5.9e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  2 TGCCACGGTACCT 15
    |||||
Db   190 TGCCACGGTACCT 177

RESULT 4
US-09-925-065A-757224/c
; Sequence 757224, Application US/09925065A
; Publication No. US2005028172A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single
; FILE REFERENCE: 108827.135
; CURRENT APPLICATION NUMBER: US/09/925,065A
; CURRENT FILING DATE: 2001-08-08
; PRIOR APPLICATION NUMBER: US 60/243,096
; PRIOR FILING DATE: 2000-10-24
; PRIOR APPLICATION NUMBER: US 60/252,147
; PRIOR FILING DATE: 2000-11-20
; PRIOR APPLICATION NUMBER: US 60/250,092
; PRIOR FILING DATE: 2000-11-30
; PRIOR APPLICATION NUMBER: US 60/261,766
; PRIOR FILING DATE: 2001-01-16
; PRIOR APPLICATION NUMBER: US 60/289,846
; PRIOR FILING DATE: 2001-05-09
; NUMBER OF SEQ ID NOS: 957086
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 757224
; LENGTH: 500
; TYPE: DNA
; ORGANISM: Homo sapiens
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US-09-925-065A-757224

Query Match      93.3%; Score 14; DB 4; Length 500;
Best Local Similarity 100.0%; Pred. No. 5.9e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  1 CTGCCACGGTACC 14
    |||||
Db   119 CTGCCACGGTACC 106

RESULT 5
US-09-738-626-586/c
; Sequence 586, Application US/09738626
; Publication No. US20020197605A1
; GENERAL INFORMATION:
; APPLICANT: NAKAGAWA, SATOSHI
; APPLICANT: MIZOGUCHI, HIROSHI
; APPLICANT: ANDO, SEIKO
; APPLICANT: HAYASHI, MIKIRO
; APPLICANT: OCHIAI, KEIKO
; APPLICANT: YOKOI, HARUHIKO
; APPLICANT: TATEISHI, NAKO
; APPLICANT: SENOH, AKIHIRO
; APPLICANT: IKEDA, MASATO
; APPLICANT: OZAKI, AKIO
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES
; FILE REFERENCE: 249-125
; CURRENT APPLICATION NUMBER: US/09/738,626
; CURRENT FILING DATE: 2000-12-18
; PRIOR APPLICATION NUMBER: JP 99/377484
; PRIOR FILING DATE: 1999-12-16
; PRIOR APPLICATION NUMBER: JP 00/159162
; PRIOR FILING DATE: 2000-04-07
; PRIOR APPLICATION NUMBER: JP 00/280988
; PRIOR FILING DATE: 2000-08-03
; NUMBER OF SEQ ID NOS: 7059
; SOFTWARE: PatentIn ver. 3.0
; SEQ ID NO 586
; LENGTH: 807
; TYPE: DNA
; ORGANISM: Corynebacterium glutamicum
US-09-738-626-586

Query Match      93.3%; Score 14; DB 3; Length 807;
Best Local Similarity 100.0%; Pred. No. 5.5e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  2 TGCCACGGTACCT 15
    |||||
Db   145 TGCCACGGTACCT 132

RESULT 6
US-10-494-675-31/c
; Sequence 31, Application US/10494675
; Publication No. US20050019877A1
; GENERAL INFORMATION:
; APPLICANT: Zelder, Oskar
; APPLICANT: Pompejus, Markus
; APPLICANT: Schroder, Hartwig
; APPLICANT: Kroger, Burkhard
; APPLICANT: Klopprogge, Corinna
; APPLICANT: Haberhauser, Gregor
; TITLE OF INVENTION: Genes coding for metabolic pathway proteins
; FILE REFERENCE: BGI-163US
; CURRENT APPLICATION NUMBER: US/10/494,675
; CURRENT FILING DATE: 2004-05-04
; PRIOR APPLICATION NUMBER: PCT/EP02/12141
; PRIOR FILING DATE: 2002-10-31
; PRIOR APPLICATION NUMBER: DE 101 54 292.1
; PRIOR FILING DATE: 2001-11-05
; NUMBER OF SEQ ID NOS: 164
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 16:59:57 ; Search time 184.56 Seconds
(without alignments)
65.779 Million cell updates/sec

Title: US-10-754-446-6

Perfect score: 15

Sequence: 1 ctgcccacggtacct 15

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 6038814 seqs, 404674181 residues

Total number of hits satisfying chosen parameters: 12077628

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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- 8: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
- 9: /cgn2_6/ptodata/2/pubpna/US11_NEW_PUB.seq.*
- 10: /cgn2_6/ptodata/2/pubpna/US60_NEW_PUB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	14	93.3	930	7	US-11-085-822-855
C 2	13.4	89.3	201	6	Sequence 855, App
C 3	13.4	89.3	708	6	Sequence 41220, A
C 4	13.4	89.3	708	6	Sequence 33670, A
C 5	13.4	89.3	847	6	Sequence 33670, A
C 6	13.4	89.3	847	6	Sequence 33670, A
C 7	13.4	89.3	1372	6	Sequence 61862, A
C 8	13.4	89.3	1372	6	Sequence 61862, A
C 9	13.4	89.3	1404	6	Sequence 57181, A
C 10	13.4	89.3	9471	6	Sequence 57181, A
C 11	13.4	89.3	168516	7	Sequence 303, App
C 12	13.4	89.3	1125000	6	Sequence 410, App
C 13	13.4	86.7	93	6	Sequence 3, Appl
C 14	13.4	86.7	441	6	Sequence 13286, A
C 15	13.4	86.7	3336	6	Sequence 17319, A
C 16	13.4	86.7	3336	6	Sequence 111, App
C 17	13.4	86.7	167116	7	Sequence 55139, A
C 18	13.4	86.7	182303	7	Sequence 55139, A
C 19	12.4	82.7	19	8	Sequence 44, Appl
C 20	12.4	82.7	19	8	Sequence 45, Appl
C 21	12.4	82.7	19	9	Sequence 51175, A
C 22	12.4	82.7	19	9	Sequence 1350721, A
C 23	12.4	82.7	20	6	Sequence 1021148, A

Sequence 1387393, A
Sequence 855540, A
Sequence 221946, A
Sequence 564915, A
Sequence 5198, Ap
Sequence 5207, Ap
Sequence 5283, Ap
Sequence 5292, Ap
Sequence 5362, Ap
Sequence 5371, Ap
Sequence 16810, A
Sequence 16812, A
Sequence 24263, A
Sequence 24481, A
Sequence 28948, A
Sequence 29049, A
Sequence 29168, A
Sequence 4350, Ap
Sequence 54379, A
Sequence 44993, A

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US-10-995-561-16810
US-10-995-561-16812
US-10-995-561-24263
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US-10-995-561-28948
US-10-995-561-29049
US-10-995-561-29168
US-11-128-061-4350
US-10-750-185-54379
US-10-750-623-54379
US-10-750-185-44993

ALIGNMENTS

RESULT 1
US-11-055-822-855/c
; Sequence 855, Application US/11055822
; Publication No. US20050260707A1
; GENERAL INFORMATION:
; APPLICANT: Pompejus, Markus
; APPLICANT: Kroger, Burkhard
; APPLICANT: Schroder, Hartwig
; APPLICANT: Zelder, Oskar
; APPLICANT: Haberhauer, Gregor
; TITLE OF INVENTION: CORNEBACTERIUM GLUTAMICUM GENES ENCODING
; FILE REFERENCE: BGI-121CPCN
; CURRENT APPLICATION NUMBER: US/11/055,822
; CURRENT FILING DATE: 2005-02-11
; PRIOR APPLICATION NUMBER: 03/606,740
; PRIOR FILING DATE: 2000-06-23
; PRIOR APPLICATION NUMBER: 60/141,031
; PRIOR FILING DATE: 1999-06-25
; PRIOR APPLICATION NUMBER: 60/142,101
; PRIOR FILING DATE: 1999-07-02
; PRIOR APPLICATION NUMBER: 60/148,613
; PRIOR FILING DATE: 1999-08-12
; PRIOR APPLICATION NUMBER: 60/187,970
; PRIOR FILING DATE: 2000-03-09
; PRIOR APPLICATION NUMBER: DE 19930476.9
; PRIOR FILING DATE: 1999-07-01
; PRIOR APPLICATION NUMBER: DE 19931415.2
; PRIOR FILING DATE: 1999-07-08
; PRIOR APPLICATION NUMBER: DE 19931418.7
; PRIOR FILING DATE: 1999-07-08
; PRIOR APPLICATION NUMBER: DE 19931419.5
; PRIOR FILING DATE: 1999-07-08
; PRIOR APPLICATION NUMBER: DE 19931420.9
; PRIOR FILING DATE: 1999-07-08
; Remaining Prior Application data removed - See File wrapper or PALM.
; NUMBER OF SEQ ID NOS: 1158
; SEQ ID NO 855
; LENGTH: 930
; TYPE: DNA
; ORGANISM: Corynebacterium glutamicum
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (101)..(907)
; OTHER INFORMATION: RXN00708
US-11-055-822-855

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Best Local Similarity 100.0%; Pred. No. 93;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB      245 TGCCACGGTACCT 232
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RESULT 2
US-10-995-561-41220/c
; Sequence 41220, Application US/10995561
; Publication No. US2005027054A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; FILE REFERENCE: DETECTION AND USES THEREOF
; FILE REFERENCE: CL001559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 41220
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-995-561-41220

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Best Local Similarity 93.3%; Pred. No. 2.1e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CTGCCACGGTACCT 15
DB      42 CTGCCACGGGACCT 28
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RESULT 3
US-10-750-185-33670/c
; Sequence 33670, Application US/10750185
; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10/750,185
; CURRENT FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 33670
; LENGTH: 708
; TYPE: DNA
; ORGANISM: Bovine 19866880906011
US-10-750-185-33670

Query Match      89.3%; Score 13.4; DB 6; Length 708;
Best Local Similarity 93.3%; Pred. No. 2e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CTGCCACGGTACCT 15
DB      106 CTGCCACGGGTACCT 92
      |||||
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RESULT 4
US-10-750-623-33670/c
; Sequence 33670, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10/750,623
; CURRENT FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 33670
; LENGTH: 708
; TYPE: DNA
; ORGANISM: Bovine 19866880906011
US-10-750-623-33670

Query Match      89.3%; Score 13.4; DB 6; Length 708;
Best Local Similarity 93.3%; Pred. No. 2e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CTGCCACGGTACCT 15
DB      106 CTGCCACGGGTACCT 92
      |||||
      |||||

RESULT 5
US-10-750-185-61862/c
; Sequence 61862, Application US/10750185
; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10/750,185
; CURRENT FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2003-12-31
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 61862
; LENGTH: 847
; TYPE: DNA
; ORGANISM: Bovine 19866881794277
US-10-750-185-61862

Query Match      89.3%; Score 13.4; DB 6; Length 847;
Best Local Similarity 93.3%; Pred. No. 2e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      1 CTGCCACGGTACCT 15
DB      177 CTGCCACGGGACCT 163
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RESULT 6
US-10-750-623-61862/c
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GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

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Run on: January 15, 2006, 16:32:47 ; Search time 54.6 Seconds
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Perfect score: 15

Sequence: 1 ctgccccacggtacct 15

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 5	13.4	89.3	273	3	US-09-668-482-33
C 6	13.4	89.3	274	3	US-08-882-164D-34
C 7	13.4	89.3	274	3	US-09-668-482-34
C 8	13.4	89.3	482	3	US-10-131-827-8312
C 9	13.4	89.3	601	3	US-09-949-016-78826
C 10	13.4	89.3	601	3	US-09-949-016-169541
C 11	13.4	89.3	601	3	US-09-949-016-169542
C 12	13.4	89.3	601	3	US-09-949-016-169543
C 13	13.4	89.3	1047	3	US-09-107-433-1277
C 14	13.4	89.3	1125	2	US-08-583-562B-9
C 15	13.4	89.3	1125	2	US-08-779-113-9
C 16	13.4	89.3	1143	3	US-09-583-110-2113
C 17	13.4	89.3	1494	2	US-08-583-562B-11
C 18	13.4	89.3	1494	2	US-08-779-113-11
C 19	13.4	89.3	1533	3	US-09-075-454-11
C 20	13.4	89.3	1660	3	US-09-347-650-11
C 21	13.4	89.3	2574	2	US-08-583-562B-1
C 22	13.4	89.3	2574	2	US-08-779-113-1
C 23	13.4	89.3	2677	3	US-08-882-164D-36
C 24	13.4	89.3	2677	3	US-09-668-482-36

C 25	13.4	89.3	3252	3	US-09-774-528-104	Sequence 104, App
C 26	13.4	89.3	3252	3	US-10-120-988-104	Sequence 104, App
C 27	13.4	89.3	3561	2	US-08-097-997A-12	Sequence 12, Appl
C 28	13.4	89.3	3561	3	US-08-665-574C-12	Sequence 12, Appl
C 29	13.4	89.3	3561	3	US-08-946-994-12	Sequence 12, Appl
C 30	13.4	89.3	4080	3	US-09-016-434-1353	Sequence 1353, Ap
C 31	13.4	89.3	4176	3	US-09-972-800A-17	Sequence 17, Appl
C 32	13.4	89.3	4176	3	US-09-023-655-1378	Sequence 1378, Ap
C 33	13.4	89.3	5238	3	US-09-620-312D-351	Sequence 351, App
C 34	13.4	89.3	7559	2	US-08-250-848-2	Sequence 2, Appl
C 35	13.4	89.3	11915	3	US-08-961-527-96	Sequence 96, Appl
C 36	13.4	89.3	12681	3	US-09-949-016-16576	Sequence 16576, A
C 37	13.4	89.3	15353	3	US-09-949-016-13454	Sequence 13454, A
C 38	13.4	89.3	17478	3	US-09-949-016-12194	Sequence 12194, A
C 39	13.4	89.3	17479	3	US-09-949-016-13057	Sequence 13057, A
C 40	13.4	89.3	44821	3	US-09-949-016-13764	Sequence 13764, A
C 41	13.4	89.3	67745	3	US-09-949-016-17251	Sequence 17251, A
C 42	13.4	89.3	110585	3	US-09-949-016-13427	Sequence 13427, A
C 43	13.4	89.3	160759	3	US-09-949-016-16514	Sequence 16514, A
C 44	13.4	89.3	784019	3	US-09-949-016-14033	Sequence 14033, A
C 45	13.4	89.3	828152	3	US-09-949-016-12777	Sequence 12777, A

ALIGNMENTS

RESULT 1

US-09-248-796A-3043

; Sequence 3043, Application US/09248796A

; Patent No. 6747137

; GENERAL INFORMATION:

; APPLICANT: Keith Weinstock et al

; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO CANDIDA ALBICAN

; TITLE OF INVENTION: FOR DIAGNOSTICS AND THERAPEUTICS

; FILE REFERENCE: 107196.132

; CURRENT APPLICATION NUMBER: US/09/248,796A

; CURRENT FILING DATE: 1999-02-12

; PRIOR APPLICATION NUMBER: US 60/074,725

; PRIOR FILING DATE: 1998-02-13

; PRIOR APPLICATION NUMBER: US 60/096,409

; PRIOR FILING DATE: 1998-08-13

; NUMBER OF SEQ ID NOS: 28208

; SEQ ID NO 3043

; LENGTH: 1323

; TYPE: DNA

; ORGANISM: Candida albicans

; US-09-248-796A-3043

Query Match 93.3%; Score 14; DB 3; Length 1323;
Best Local Similarity 100.0%; Pred. No. 2.8e+02;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTGCCACGGTACC 14

Db 1025 CTGCCACGGTACC 1038

RESULT 2

US-09-221-017B-510

; Sequence 510, Application US/09221017B

; Patent No. 6444799

; GENERAL INFORMATION:

; APPLICANT: Ross, Bruce C.

; TITLE OF INVENTION: P. GINGIVALIS NUCLEOTIDES AND USES THEREOF

; NUMBER OF SEQUENCES: 1120

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: MORRISON & FOERSTER

; STREET: 755 PAGE MILL ROAD

; CITY: Palo Alto

; STATE: CA

; COUNTRY: USA

; ZIP: 94304-1018

; COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FastSO for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/221,017B
FILING DATE: 23-DEC-1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P11182
FILING DATE: 31-DEC-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P11546
FILING DATE: 30-JAN-1998
PRIOR APPLICATION DATA:
APPLICATION NUMBER: P22911
FILING DATE: 09-APR-1998
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/AU98/01023
FILING DATE: 10-DEC-1998
ATTORNEY/AGENT INFORMATION:
NAME: MOROY, Gladys H
REGISTRATION NUMBER: 32,430
REFERENCE/DOCKET NUMBER: 27340-20021.00
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-813-5600
TELEFAX: 650-494-0792
TELEX: 706141
INFORMATION FOR SEQ ID NO: 510:
SEQUENCE CHARACTERISTICS:
LENGTH: 7319 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: circular
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: UNKNOWN
ORIGINAL SOURCE:
ORGANISM: PORPHYROMONAS GINGIVALIS
FEATURE:
NAME/KEY: misc_feature
LOCATION: 1...7319
US-09-221-017B-510

Query Match 93.3%; Score 14; DB 3; Length 7319;
Best Local Similarity 100.0%; Pred. No. 2.9e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 3741 CTGCCCACGGTACC 3754

RESULT 3
US-08-707-237A-5
Sequence 5, Application US/08707237A
Patent No. 5830713
GENERAL INFORMATION:
APPLICANT: Ferrari, Franco A.
APPLICANT: Capello, Joseph
APPLICANT: Crissman, John W.
APPLICANT: Dorman, Mary A.
TITLE OF INVENTION: METHODS FOR PREPARING SYNTHETIC
TITLE OF INVENTION: REPETITIVE DNA
NUMBER OF SEQUENCES: 108
CORRESPONDENCE ADDRESS:
ADDRESSEE: Flehr, Hobbach, Test, Albritton & Herbert
STREET: Four Embarcadero Center, Suite 3400
CITY: San Francisco
STATE: California
COUNTRY: United States
ZIP: 94111-4187
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/707,237A
FILING DATE: 03-SEP-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/175,155
FILING DATE: 29-DEC-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/053,049
FILING DATE: 22-APR-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/609,716
FILING DATE: 06-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/269,429
FILING DATE: 09-NOV-1988
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/114,618
FILING DATE: 29-OCT-1987
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 06/927,258
FILING DATE: 04-NOV-1986
ATTORNEY/AGENT INFORMATION:
NAME: Trecartin, Richard P.
REGISTRATION NUMBER: 31,801
REFERENCE/DOCKET NUMBER: A-55186-10/WHO
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 781-1989
TELEFAX: (415) 398-3249
TELEX: 910 277299
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 63 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
US-08-707-237A-5

Query Match 89.3%; Score 13.4; DB 2; Length 63;
Best Local Similarity 93.3%; Pred. No. 5.8e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 26 CTACCCACGGTACCT 40

RESULT 4
US-08-882-164D-33/c
Sequence 33, Application US/08882164D
Patent No. 6306624
GENERAL INFORMATION:
APPLICANT: Petkovich, P. Martin, White, Jay A.,
APPLICANT: Beckett, Barbara R., Jones, Glenville
TITLE OF INVENTION: Retinoid Metabolizing Protein
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: Blake, Casaeals & Graydon
STREET: Box 25, Commerce Court West
CITY: Toronto
STATE: Ontario
COUNTRY: Canada
ZIP: M5L 1A9
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3 1/2 inch, 1.4 Mb storage
COMPUTER: COMPAQ, IBM PC compatible
OPERATING SYSTEM: MS-DOS 5.1
SOFTWARE: WORD PERFECT

GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 785.792 Seconds
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Title: US-10-754-446-7

Perfect score: 16

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Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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C 4	16	100.0	661	10	G93242 S208P6433RE
5	16	100.0	728	10	BV524331 G591P6065
C 6	16	100.0	925	10	BV526258 G591P6047
7	16	100.0	1740	6	AX280021 Sequence
8	16	100.0	1829	6	CQ719994 Sequence
9	16	100.0	2004	8	AF249319
10	16	100.0	2037	8	HSA293970
11	16	100.0	2049	8	AK026102
12	16	100.0	2051	8	AF287269
13	16	100.0	2052	8	AX083508
14	16	100.0	2063	8	AK222673
15	16	100.0	2065	8	AB125179
16	16	100.0	2078	8	BC005149
17	16	100.0	2094	6	BD233734
18	16	100.0	2095	6	AX280019 Sequence

19	16	100.0	2272	8	HSA2933659
C 20	16	100.0	12300	1	AE005887
21	16	100.0	13270	8	AF287270
22	16	100.0	47196	14	BX546462.4
C 23	16	100.0	49264	14	AC164672
C 24	16	100.0	128501	8	AC012614
C 25	16	100.0	150909	9	AC116811
26	16	100.0	151130	9	AC127432
C 27	16	100.0	155645	14	AC021153
28	16	100.0	157952	8	AF003691
C 29	16	100.0	158063	8	AP001046
30	16	100.0	165994	9	AC129207
31	16	100.0	173081	9	AC159338
32	16	100.0	173126	8	AC086878
C 33	16	100.0	175177	9	AC124383
34	16	100.0	179556	14	AC018734
35	16	100.0	180770	14	AC153861
36	16	100.0	188593	8	AC016877
37	16	100.0	194042	9	AC153850
C 38	16	100.0	194364	14	AC164574
C 39	16	100.0	208898	14	AC164628
C 40	16	100.0	210408	8	AC010608
41	16	100.0	213826	9	AL845298
42	16	100.0	223368	14	AC095138
C 43	16	100.0	225335	14	AC111381
C 44	16	100.0	232747	14	AC116259
45	16	100.0	253693	14	AC131839

ALIGNMENTS

RESULT 1
BV401493/c

LOCUS

DEFINITION S229P6332R8.T0 GermanShepherd Canis familiaris STS genomic,
sequence tagged site.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BV401493

S229P6332R8.T0 GermanShepherd Canis familiaris STS genomic,

sequence tagged site.

BV401493

SV401493.1 GI:57775274

STS.

Canis familiaris (dog)

Canis familiaris

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Laurasiatheria; Carnivora; Fissipedia; Canidae;

Canis.

1 (bases 1 to 536)

Lindblad-Toh, K.

The genome sequence of Canis familiaris

Unpublished (2004)

Contact: Kerstin Lindblad-Toh

Whitehead Institute for Biomedical Research, Center for Genome

Research

320 Charles Street, Cambridge, MA 02141, USA

Tel: 6172521477

Fax: 6172580903

Email: kersli@genome.wi.mit.edu

Primer A: No sequence submitted

Primer B: No sequence submitted

STS size: 536

Protocol:

WGS-discovery (WGS):

Paired-end low-coverage whole genome shotgun reads were generated

from 9 breeds

(German Shepherd, Rottweiler, Bedlington Terrier, Beagle, Labrador

Retriever, English

Shepherd, Italian Greyhound, Alaskan Malamute and the Portuguese

Water Dog -100,000 each)

and five other canids (Chinese, Alaskan, Indian and Spanish Gray

Wolf as well as the

Californian Coyote).

The WGS reads were placed uniquely on the CanFam1.0 boxer assembly

and SNP detection was

carried out by SSAHA-SNP. 863872 reads were annotated as STSs and 485941 SNPs were annotated with alleles from the boxer and the breed or canid from which the particular read came. The validation rate for these SNPs was estimated at approximately 98%.

WGA-discovery (WGA) of Boxer/Poodle SNPs: A second set of SNPs was generated using a similar methodology except that the contigs from the 1.5x poodle assembly (Kirkness 2003) were used instead of WGS reads. Since this sequence lacked base quality scores, arbitrary quality scores of phred 40 were assigned before the poodle sequence was placed uniquely on the CanFam1.0 boxer assembly and SNP detection was carried out by SSAHA-SNP. 1637780 SNPs were annotated with alleles from the boxer and the breed or canid from which the particular read came. The validation rate for these SNPs was estimated at approximately 98%.

Internal-WGA-discovery (I-WGA): A third set of SNPs were discovered by comparing reads in the WGA assembly. SNPs were defined as mismatch positions that had a base quality of ≥ 30 on both reads in a region that aligned without gaps, and with at most one additional mismatch in the ten flanking bases. For each allele, at least one additional read had to confirm it. 731476 SNPs were annotated with alleles between the two boxer alleles. The validation rate for these SNPs was estimated at approximately 98%.

FEATURES
source
1. .536
/organism="Canis familiaris"
/mol_type="genomic DNA"
/strain="GermanShepherd"
/db_xref="taxon:9615"
/map="8 22-462 74954487-74954047"
/clone_lib="GermanShepherd"
1. .536

STS
ORIGIN

Query Match 100.0%; Score 16; DB 10; Length 536;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACCCAGGCCACAT 16
|||||
Db 468 AGACCCAGGCCACAT 453

RESULT 2
AF305572S5
LOCUS AF305572S5 579 bp DNA linear PRI 26-DEC-2000
DEFINITION Homo sapiens mucopolipin 1 (MCOLN1) gene, exons 6 and 7.
ACCESSION AF305576
VERSION AF305576.1 GI:11991202
KEYWORDS
SEGMENT 5 of 8
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1 (bases 1 to 579)
Bargal, R., Avidan, N., Ben-Asher, E., Olender, Z., Zeigler, M.,
Frumkin, A., Raas-Rothschild, A., Glusman, G., Lancet, D., and Bach, G.
Identification of the gene causing mucopolipidosis type IV
Nat. Genet. 26 (1), 118-123 (2000)
10973263
REFERENCE 2 (bases 1 to 579)
Bargal, R., Avidan, N., Ben-Asher, E., Olender, A., Zeigler, M.,
Frumkin, A., Raas-Rothschild, A., Glusman, G., Lancet, D., and Bach, G.

TITLE
JOURNAL
FEATURES
source
1. .579
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
129_225
/gene="MCOLN1"
/number=6
423. .522
/gene="MCOLN1"
/number=7

ORIGIN

Query Match 100.0%; Score 16; DB 8; Length 579;
Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACCCAGGCCACAT 16
|||||
Db 472 AGACCCAGGCCACAT 487

RESULT 3
BV230130/c
LOCUS BV230130 659 bp DNA linear STS 19-JAN-2005
DEFINITION S233P6354FD9.T0 LabradorRetriever Canis familiaris STS genomic,
sequence tagged site.
ACCESSION BV230130
VERSION BV230130.1 GI:57292672
KEYWORDS STS.
SOURCE Canis familiaris (dog)
ORGANISM Canis familiaris
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Laurasiatheria; Carnivora; Fissipedia; Canidae;
Canis.
REFERENCE 1 (bases 1 to 659)
Lindblad-Toh, K.
AUTHORS The genome sequence of Canis familiaris
TITLE Unpublished (2004)
JOURNAL
COMMENT
Contact: Kerstin Lindblad-Toh
Whitehead Institute for Biomedical Research, Center for Genome
Research
320 Charles Street, Cambridge, MA 02141, USA
Tel: 6172521477
Fax: 6172580903
Email: kersli@genome.wi.mit.edu
Primer A: No sequence submitted
Primer B: No sequence submitted
STS size: 659
Protocol:
WGS-discovery (WGS):
Paired-end low-coverage whole genome shotgun reads were generated
from 9 breeds
(German Shepherd, Rottweiler, Bedlington Terrier, Beagle, Labrador
Retriever, English
Shepherd, Italian Greyhound, Alaskan Malamute and the Portuguese
Water Dog -100,000 each)
and five other canids (Chinese, Alaskan, Indian and Spanish Gray
Wolf as well as the
Californian Coyote).
The WGS reads were placed uniquely on the CanFam1.0 boxer assembly
and SNP detection was
carried out by SSAHA-SNP. 863872 reads were annotated as STSs and
485941 SNPs were
annotated with alleles from the boxer and the breed or canid from
which the particular
read came. The validation rate for these SNPs was estimated at
approximately 98%.

WGA-discovery (WGA) of Boxer/Poodle SNPs:

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 1789.31 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-7
Perfect score: 16
Sequence: 1 agaccaggccacat 16

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*
1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_hc.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_est7.*
9: gb_gss1.*
10: gb_gss2.*
11: gb_gss3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	100.0	278	9	AZ491094
2	16	100.0	291	8	T29948
3	16	100.0	316	8	F07345
4	16	100.0	324	1	AW486159
5	16	100.0	331	8	Z43611
6	16	100.0	333	8	T35364
7	16	100.0	350	8	Z43604
8	16	100.0	371	1	AW356454
9	16	100.0	387	1	AW425724
10	16	100.0	387	2	B5478501
11	16	100.0	404	9	AZ847535
12	16	100.0	442	1	AW785343
13	16	100.0	466	9	AZ073212
14	16	100.0	478	2	BG383183
15	16	100.0	500	2	BF443224
16	16	100.0	510	6	CA889269
17	16	100.0	540	7	CJ017244
18	16	100.0	578	9	AZ286475
19	16	100.0	593	7	CN788730
20	16	100.0	740	6	CX489568
21	16	100.0	772	7	CK465307
22	16	100.0	775	10	CG740895

c	23	16	100.0	777	7	CK467406
c	24	16	100.0	779	7	CK468501
c	25	16	100.0	817	2	B1252802
c	26	16	100.0	826	2	BG913075
c	27	16	100.0	845	2	BG176074
c	28	16	100.0	862	7	CK469557
c	29	16	100.0	868	3	B1909885
c	30	16	100.0	872	10	CZ363216
c	31	16	100.0	873	6	CD558014
c	32	16	100.0	878	5	BQ956452
c	33	16	100.0	901	5	BX433310
c	34	16	100.0	961	10	CZ363668
c	35	16	100.0	979	5	BQ918296
c	36	16	100.0	993	2	BG739714
c	37	16	100.0	1056	11	CNS037W4
c	38	16	100.0	1078	5	BX362786
c	39	16	100.0	1273	4	CR616477
c	40	16	100.0	1712	10	AY410284
c	41	16	100.0	1743	10	AY410283
c	42	16	100.0	1996	4	CR622331
c	43	16	100.0	2108	4	AK052519
c	44	15.6	97.5	961	5	BX397924
c	45	15.2	95.0	1026	5	BX364018

ALIGNMENTS

RESULT 1
AZ491094
LOCUS
DEFINITION
IM0324P22F Mouse 10kb plasmid UUGCLM library Mus musculus genomic clone UUGCLM0324P22 F, genomic survey sequence.
ACCESSION
AZ491094
VERSION
AZ491094.1 GI:10662462
KEYWORDS
GSS.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muroidea; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 278)
Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A. and Wright, D., Weiss, R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
Unpublished (2000)
JOURNAL
COMMENT
Contact: Robert B. Weiss
University of Utah Genome Center
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0324 row: P column: 22
Seq primer: CGTTGTAACACGCGCCAGT
Class: plasmid ends
High quality sequence stop: 278.
Location/Qualifiers
1. .278
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGCLM0324P22"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/clone_lib="Mouse 10kb plasmid library"
/note="Vector: PWD42hv; Purified genomic DNA from M.

FEATURES
source

1. .278
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGCLM0324P22"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/clone_lib="Mouse 10kb plasmid library"
/note="Vector: PWD42hv; Purified genomic DNA from M.

musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gii4732114[gb|AF129072.1]), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent *E. coli* XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 100.0%; Score 16; DB 9; Length 278;
Best Local Similarity 100.0%; Pred. NO. 4e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
|||||
Db 129 AGACCCAGGCCACAT 144

RESULT 2

T29948
LOCUS T29948.1 GI:612046
DEFINITION EST100452 Human Pancreas Homo sapiens cDNA 5' end similar to None, mRNA sequence.

ACCESSION T29948.1
VERSION GI:612046
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 291)
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A., Bult,C.J., Lee,N., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D., White,O., Sutton,G., Blake,J.A., Brandon,R.C., Chiu,M.-W., Clayton,R.A., Cline,R.T., Cotton,M.D., Earle-Hughes,J., Fine,L.D., FitzGerald,L.M., FitzHugh,W.M., Fritchman,J.L., Geoghagen,N.S.M., Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., HinkleJr,P.S., Kelley,J.M., Kline,K.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M., Moreno-Balances,R.F., McDonald,L.A., Nguyen,D.T., Pellegrino,S.M., Phillips,C.A., Ryder,S.E., Scott,J.L., Sauddek,D.M., Shirley,R., Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y., Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.-J., Dimke,D., Feng,P., Ferrie,A., Fischer,C., Hastings,G.A., He,W.-W., Hu,J.-S., Greene,J.M., Gruber,J., Hudson,P., Kim,A., Kozak,D.L., Kunsch,C., Ji,H., Li,H., Meisner,P.S., Olsen,H., Raymond,L., Wei,Y.-F., Wing,J., Xu,C., Yu,G.-L., Ruben,S.M., Dillon,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C., Fraser,C.M. and Venter,J.C.
Initial Assessment of Human Gene Diversity and Expression Patterns Based Upon 83 Million Basepairs of cDNA Sequence
Nature 377, 3-174 (1995)

TITLE
JOURNAL Nature 377, 3-174 (1995)
PUBMED 7566098

COMMENT

Other_ESTs: THC11326
Contact: Venter, JC
The Institute for Genomic Research
932 Clopper Rd, Gaithersburg, MD 20878
Tel: 3018698056
Fax: 3018699423
Email: tdbinfo@tdb.tigr.org
For clone availability, additional sequence and expression information related to this EST, please contact the TIGR Database

(tdbinfo@tdb.tigr.org)
Seq primer: M13 Reverse.
Location/Qualifiers
1. .291
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="ATCC (inhost):107961"
/db_xref="taxon:9606"
/clone_lib="Human Pancreas"
/note="Organ: pancreas"

ORIGIN

Query Match 100.0%; Score 16; DB 8; Length 291;
Best Local Similarity 100.0%; Pred. No. 4e+03;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
|||||
Db 186 AGACCCAGGCCACAT 201

RESULT 3

F07345
LOCUS F07345
DEFINITION HSC22G041 normalized infant brain cDNA Homo sapiens CDNA clone c-22g04, mRNA sequence.

ACCESSION F07345.1
VERSION GI:672999
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 316)
Auffray,C., Behar,G., Bois,F., Bouchier,C., da Silva,C., Devignes,M.D., Duprat,S., Houlgatte,R., Jumeau,M.N., Lamy,B., Lorenzo,F., Mitchell,H., Mariage-Samson,R., Pietu,G., Pouliot,Y., Sebastiani-Kabaktchis,C. and Tessier,A.
IMAGE: molecular integration of the analysis of the human genome and its expression
C. R. Acad. Sci. III, Sci. Vie 318 (2), 263-272 (1995)
7757816

COMMENT

Contact: Genethon
Genexpress-Genethon
Genethon Centre de recherche sur le Genome Humain
1,rue de l'Internationale, BP60 91002 EVRY Cedex, FRANCE
Tel: 33169472800
Fax: 33160778698
Email: genexpress@genethon.fr
Single read
Genexpress_library_idt: C; Genexpress_sequence_idt: ylc-22g04
Seq primer: (-21)M13 universal.

FEATURES

source

1. .316
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="c-22g04"
/sex="Female"
/tissue_type="total brain"
/dev_stage="3 months old"
/clone_lib="normalized infant brain cDNA"
/notes="Organ: brain; Vector: lafmid BA; Site_1: HindIII; Site_2: NotI; sex=Female; dev_stage=3 months old; isolate=muscular atrophy patient; tissue_type=total brain; total mRNA was oligo-(dT) primed and directionally cloned 5' -> 3' into the HindIII -> NotI sites of the lafmid BA vector. Clone library from B. Soares, Psychiatry Dept. Columbia University, USA. Normalization_method: Bento Soares, P.N.A.S in press"

ORIGIN

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 217.088 Seconds
(without alignments)
491.207 Million cell updates/sec

Title: US-10-754-446-7

Perfect score: 16

Sequence: 1 agaccaggccacat 16

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

- N_Geneseq_21.*
1: Geneseqn1980s.*
2: Geneseqn1990s.*
3: Geneseqn2000s.*
4: Geneseqn2001as.*
5: Geneseqn2001bs.*
6: Geneseqn2002as.*
7: Geneseqn2002bs.*
8: Geneseqn2003as.*
9: Geneseqn2003bs.*
10: Geneseqn2003cs.*
11: Geneseqn2003ds.*
12: Geneseqn2004as.*
13: Geneseqn2004bs.*
14: Geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16	100.0	16	14 AEB28963	Aeb28963 Human MCO
2	16	100.0	1619	4 AAI61022	Aai61022 Human pol
3	16	100.0	1740	6 AAI171700	Aai171700 Human TRP
4	16	100.0	1740	6 ABL40755	Abi40755 Human TLC
5	16	100.0	1741	4 AAI59236	Aai59236 Human pol
6	16	100.0	1743	9 ADB84284	Adb84284 Human nuc
7	16	100.0	1743	14 ADV66231	Adv66231 TRP-like
8	16	100.0	2051	13 ADP25150	Adp25150 PRO polyp
9	16	100.0	2052	4 AAF81753	Aaf81753 Human mem
10	16	100.0	2092	5 AAS72274	Aas72274 DNA encod
11	16	100.0	2092	6 ABL90358	Abi90358 Human pol
12	16	100.0	2094	3 AAA39067	Aaa39067 Human sec
13	16	100.0	2095	6 AAI171699	Aai171699 Human TRP
14	16	100.0	2095	6 ABL40754	Abi40754 Human TLC
15	16	100.0	2095	14 ADV66229	Adv66229 TRP-like
16	16	100.0	2140	12 ADQ24925	Adq24925 Human sof
17	16	100.0	11365	4 AAK73827	Aak73827 Human imm
18	16	100.0	13270	9 ADB84283	Adb84283 Human nuc
19	16	100.0	20046	4 AAK73826	Aak73826 Human imm

20	16	100.0	340449	8 AAL52198	Aal52198 Human sec
c 21	15	93.8	384	6 ABNI5902	Abni5902 Human ORF
c 22	15	93.8	449	11 ADT97636	Adt97636 Colon can
c 23	15	93.8	449	11 ADX44118	Adx44118 Human CDN
c 24	15	93.8	471	13 ADU10604	Adu10604 Solid tum
c 25	15	93.8	502	4 AAK61151	Aak61151 Human imm
c 26	15	93.8	514	13 ADQ49858	Adq49858 Novel can
c 27	15	93.8	625	4 AAH05091	Aah05091 Human CDN
c 28	15	93.8	819	3 AAC79832	Aac79832 Human sec
c 29	15	93.8	882	4 AAH34308	Aah34308 Human col
c 30	15	93.8	882	6 ABL90324	Abi90324 Human pol
c 31	15	93.8	943	4 AAK79476	Aak79476 Human imm
c 32	15	93.8	943	4 AAK79477	Aak79477 Human imm
c 33	15	93.8	1422	5 AAS83077	Aas83077 DNA encod
c 34	15	93.8	1468	4 AAH15702	Aah15702 Human CDN
c 35	15	93.8	1957	10 ADE59228	Ades59228 Human gen
c 36	15	93.8	1957	10 ADE59225	Ades59225 Human gen
c 37	15	93.8	2000	11 ACL37128	Acl37128 Rice stre
c 38	15	93.8	2004	10 ADB63068	Adb63068 Human CDN
c 39	15	93.8	3111	12 ADN38521	Adn38521 Novel hum
c 40	15	93.8	3444	11 ADL22622	Adl22622 Human dis
c 41	15	93.8	3600	8 ADA83759	Ada83759 Human KIA
c 42	15	93.8	3600	12 ADI24494	Adi24494 Human mod
c 43	15	93.8	3600	14 ADX83158	Adx83158 Human TEG
c 44	15	93.8	3600	14 ADX44432	Adx44432 Human col
c 45	15	93.8	3600	14 ADY39343	Ady39343 Human col

ALIGNMENTS

RESULT 1

AEb28963	ID	AEb28963 standard; DNA; 16 BP.
XX	AC	AEb28963;
XX	DT	22-SEP-2005 (first entry)
XX	DE	Human MCOLN1 gene probe DEL SEQ ID NO:7.
XX	KW	mcucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; probe;
XX	OS	ss.
XX	PN	Homo sapiens.
XX	PD	US2005153300-A1.
XX	PF	14-JUL-2005.
XX	PR	09-JAN-2004; 2004US-00754446.
XX	PR	09-JAN-2004; 2004US-00754446.
XX	PR	(QUES-) QUEST DIAGNOSTICS INC.
XX	PI	Sun W, Rantash F;
XX	DR	WPI; 2005-521160/53.

Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by amplifying the nucleic acid, detecting amplified product with labeled oligonucleotide probes via a change in fluorescence which indicates the presence of an ML IV mutant.

Claim 3; SEQ ID NO 7; 15pp; English.

The invention relates to a method (M1) for detecting the presence of mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1) involves contacting the nucleic acid with oligonucleotide primers and probes, conducting amplification by temperature cycling and monitoring the accumulation of amplified nucleic acid by detecting an increase in donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucolin-1
 CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a probe for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

XX
 SQ Sequence 16 BP; 5 A; 7 C; 3 G; 1 T; 0 U; 0 Other;
 Query Match 100.0%; Score 16; DB 14; Length 16;
 Best Local Similarity 100.0%; Pred. No. 3.8e+02; Mismatches 0; Indels 0; Gaps 0;
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QY 1 AGACCCAGGCCACAT 16
 Db 1 AGACCCAGGCCACAT 16
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RESULT 2
 AAI61022/c
 ID AAI61022 standard; cDNA; 1619 BP.
 AC AAI61022;
 XX
 DT 22-OCT-2001 (first entry)
 XX
 DE Human polynucleotide SEQ ID NO 5011.
 XX
 KW Human; neurotropic; immunosuppressant; cytostatic; gene therapy; cancer;
 KW peripheral nervous system; neuropathy; central nervous system; CNS;
 KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
 KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
 KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
 KW leukaemia; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200153312-A1.
 XX
 PD 26-JUL-2001.
 XX
 PF 26-DEC-2000; 2000WO-US034263.
 XX
 PR 23-DEC-1999; 99US-00471275.
 PR 21-JAN-2000; 2000US-00488725.
 PR 25-APR-2000; 2000US-00552317.
 PR 20-JUN-2000; 2000US-00598042.
 PR 19-JUL-2000; 2000US-00620312.
 PR 03-AUG-2000; 2000US-00653450.
 PR 14-SEP-2000; 2000US-00662191.
 PR 19-OCT-2000; 2000US-00693036.
 PR 29-NOV-2000; 2000US-00727344.
 XX
 PA (HYSE-) HYSEQ INC.
 XX
 PI Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
 PI Wang J, Wang J, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J, Zhao QA;
 PI Zhou P, Goodrich R, Drmanac RT;
 XX
 DR WPI; 2001-442253/47.
 DR P-PSDB; AAM41866.
 XX
 XX Novel nucleic acids and polypeptides, useful for treating disorders such
 PT as central nervous system injuries.
 PT
 XX
 PS Claim 1; SEQ ID NO 5011; 10078pp; English.
 XX
 CC The invention relates to human nucleic acids (AAI57798-AAI61369) and the

CC encoded polypeptides (AAM38642-AAM42213) with neurotropic,
 CC immunosuppressant and cytostatic activity. The polynucleotides are useful
 CC in gene therapy. A composition containing a polypeptide or polynucleotide
 CC of the invention may be used to treat diseases of the peripheral nervous
 CC system, such as peripheral nervous injuries, peripheral neuropathy and
 CC localised neuropathies and central nervous system diseases, such as
 CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
 CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
 CC utilisation of the activities such as: Immune system suppression,
 CC activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
 CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
 CC assays for receptor activity, arthritis and inflammation, leukaemia and
 CC C.N.S disorders. Note: The sequence data for this patent did not form
 CC part of the printed specification

XX
 SQ Sequence 1619 BP; 350 A; 450 C; 518 G; 301 T; 0 U; 0 Other;
 Query Match 100.0%; Score 16; DB 4; Length 1619;
 Best Local Similarity 100.0%; Pred. No. 4.3e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
 Db 1122 AGACCCAGGCCACAT 1107
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RESULT 3
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 ID AAI71700 standard; cDNA; 1740 BP.
 XX
 AC AAI71700;
 XX
 DT 29-JAN-2002 (first entry)
 XX
 DE Human TRP-like calcium channel TLCC-2 coding sequence #2.
 XX
 KW Human; TLCC-2; TRP-like calcium channel; membrane excitability;
 KW nociception; neurotropic; neuroprotective; antiparkinsonian; cytostatic;
 KW hypotensive; antidepressant; analgesic; anticonvulsant; tranquiliser;
 KW Parkinson's disease; Huntington's disease; multiple sclerosis;
 KW Gilles de la Tourette's syndrome; autonomic function disorder; cancer;
 KW neuroleptic; gene therapy; Alzheimer's disease; CNS disorder; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT CDS 1..1740
 FT /*tag= a
 FT /product= "TLCC-2"
 FT /partial
 XX
 PN WO200177331-A1.
 XX
 PD 18-OCT-2001.
 XX
 PF 06-APR-2001; 2001WO-US011442.
 XX
 PR 07-APR-2000; 2000US-00544797.
 XX
 PA (MILL-) MILLENIUM PHARM INC.
 XX
 PI Curtis RAJ, Silos-Santiago I;
 XX
 DR WPI; 2002-010913/01.
 DR P-PSDB; AAM51858.
 XX
 XX Novel isolated human transient receptor potential-like calcium channel
 PT protein-2 useful for treating Alzheimer's disease, depression, amnesia,
 PT pain disorder, and cancer.
 PT
 XX
 PS Claim 1; Fig 1; 148pp; English.
 XX
 CC The present invention relates to the protein and coding sequences of

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 16:41:33 ; Search time 386.432 Seconds
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342.389 Million cell updates/sec

Title: US-10-754-446-7

Perfect score: 16

Sequence: 1 agaccaggccacat 16

Scoring table: IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 9793542 seqs, 4134689005 residues

Total number of hits satisfying chosen parameters: 19587084

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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9: /cgn2_6/ptodata/1/pubpna/US10E_PUBCOMB.seq.*
10: /cgn2_6/ptodata/1/pubpna/US11_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	16	100.0	1400	10 US-11-060-756-2850	Sequence 2850, Ap
3	16	100.0	1400	10 US-11-060-756-7122	Sequence 7122, Ap
4	16	100.0	1740	3 US-09-828-466-3	Sequence 3, Appli
5	16	100.0	1740	5 US-10-103-458-3	Sequence 3, Appli
6	16	100.0	1743	8 US-10-782-695-6	Sequence 6, Appli
7	16	100.0	2051	3 US-09-851-494B-2	Sequence 2, Appli
8	16	100.0	2052	3 US-09-965-529-50	Sequence 50, Appli
9	16	100.0	2052	3 US-09-969-680A-50	Sequence 50, Appli
10	16	100.0	2052	10 US-11-048-692-50	Sequence 920, App
11	16	100.0	2092	6 US-10-284-237-920	Sequence 920, App
12	16	100.0	2092	9 US-10-450-763-8078	Sequence 8078, Ap
13	16	100.0	2094	3 US-09-820-893-26	Sequence 26, Appl
14	16	100.0	2094	7 US-10-607-565-26	Sequence 26, Appl
15	16	100.0	2095	3 US-09-828-466-1	Sequence 1, Appli
16	16	100.0	2095	5 US-10-103-458-1	Sequence 1, Appli
17	16	100.0	2095	8 US-10-782-695-4	Sequence 4, Appli
18	16	100.0	2140	8 US-10-723-860-7745	Sequence 7745, Ap
19	16	100.0	13270	3 US-09-851-494B-1	Sequence 1, Appli
20	16	100.0	34049	3 US-09-903-582-3	Sequence 3, Appli
21	15	93.8	342	7 US-10-437-963-97094	Sequence 97094, A
22	15	93.8	445	5 US-10-086-543-3155	Sequence 3155, Ap
23	15	93.8	552	7 US-10-767-701-22677	Sequence 22677, A

Sequence 531750,
Sequence 220987,
Sequence 220987,
Sequence 502281,
Sequence 73411, A
Sequence 73412, A
Sequence 73413, A
Sequence 647660,
Sequence 703658,
Sequence 703659,
Sequence 703660,
Sequence 77806, A
Sequence 1400, Ap
Sequence 886, App
Sequence 18881, A
Sequence 1222, Ap
Sequence 41, Appl
Sequence 52, Appl
Sequence 52, Appl
Sequence 1006, Ap
Sequence 116, App
Sequence 853, App

24 15 93.8 558 4 US-09-925-065A-531750
c 25 15 93.8 596 5 US-10-027-632-220987
c 26 15 93.8 596 6 US-10-027-632-220987
c 27 15 93.8 609 4 US-09-925-065A-502281
c 28 15 93.8 614 4 US-09-925-065A-73411
c 29 15 93.8 614 4 US-09-925-065A-73412
c 30 15 93.8 614 4 US-09-925-065A-73413
c 31 15 93.8 627 4 US-09-925-065A-647660
c 32 15 93.8 675 4 US-09-925-065A-703658
c 33 15 93.8 675 4 US-09-925-065A-703659
c 34 15 93.8 675 4 US-09-925-065A-703660
c 35 15 93.8 678 8 US-10-425-115-77806
c 36 15 93.8 882 5 US-10-106-698-1400
c 37 15 93.8 882 6 US-10-284-237-886
c 38 15 93.8 1422 9 US-10-450-763-18881
c 39 15 93.8 2004 6 US-10-104-047-1222
c 40 15 93.8 3600 5 US-10-157-031-41
c 41 15 93.8 3600 8 US-10-651-237-52
c 42 15 93.8 3600 8 US-10-782-413-52
c 43 15 93.8 52510 5 US-10-087-192-1006
c 44 15 93.8 59747 7 US-10-450-826-116
c 45 15 93.8 197775 5 US-10-087-192-853

ALIGNMENTS

RESULT 1
US-10-754-446-7
; Sequence 7, Application US/10754446
; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: SUN, WEIMIN
; APPLICANT: HANTASH, FERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; TITLE OF INVENTION: MUCOLIPIDOSIS IV MUTATIONS
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 3.2
; SEQ ID NO 7
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Probe
US-10-754-446-7

Query Match 100.0%; Score 16; DB 9; Length 16;
Best Local Similarity 100.0%; Pred. No. 4.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACCCAGGCCACAT 16
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Db 1 AGACCCAGGCCACAT 16

RESULT 2
US-11-060-756-2850
; Sequence 2850, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; TITLE OF INVENTION: Target Genes
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2850

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; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-2850

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Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 301 AGACCCAGGCCACAT 316

RESULT 3
US-11-060-756-7122
; Sequence 7122, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; TARGET GENES:
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7122
; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-7122

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Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
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Db 301 AGACCCAGGCCACAT 316

RESULT 4
US-09-828-466-3
; Sequence 3, Application US/09828466
; Patent No. US20020035056A1
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Silos-Santiago, Immaculada
; TITLE OF INVENTION: 54420, A NOVEL HUMAN CALCIUM CHANNEL
; FILE REFERENCE: MNI-125CP
; CURRENT APPLICATION NUMBER: US/09/828,466
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 09/544,797
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 1740
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1740)
US-09-828-466-3

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Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 15; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
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Db 827 AGACCCAGGCCACAT 842

RESULT 5
US-10-103-458-3
; Sequence 3, Application US/10103458
; Publication No. US20020197680A1
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; TITLE OF INVENTION: 54420, A NOVEL HUMAN CALCIUM CHANNEL
; FILE REFERENCE: MNI-125
; CURRENT APPLICATION NUMBER: US/10/103,458
; CURRENT FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: US/09/544,797
; PRIOR FILING DATE: PEOE FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 3
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US-10-103-458-3

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Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
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Db 827 AGACCCAGGCCACAT 842

RESULT 6
US-10-782-695-6
; Sequence 6, Application US/10782695
; Publication No. US20040248160A1
; GENERAL INFORMATION:
; APPLICANT: Glucksmann, Maria A.
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Tsai, Fong-Ying
; APPLICANT: Hodge, Martin R.
; APPLICANT: Meyers, Rachel E.
; APPLICANT: MacBeth, Kyle J.
; APPLICANT: Bandaru, Rajasekhar
; TITLE OF INVENTION: NOVEL 14275, 54420, 8797, 27439, 68730,
; FILE REFERENCE: 69112 AND 52908 MOLECULES AND USES THEREFOR
; CURRENT APPLICATION NUMBER: US/10/782,695
; CURRENT FILING DATE: 2004-02-19
; PRIOR APPLICATION NUMBER: US 09/945,254
; PRIOR FILING DATE: 2001-08-31
; PRIOR APPLICATION NUMBER: US 60/229,829
; PRIOR FILING DATE: 2000-08-31
; PRIOR APPLICATION NUMBER: US 09/945,301
; PRIOR FILING DATE: 2001-08-31
; PRIOR APPLICATION NUMBER: US 60/229,301
; PRIOR FILING DATE: 2000-09-01
; PRIOR APPLICATION NUMBER: US 10/007,399
; PRIOR FILING DATE: 2001-11-05
; PRIOR APPLICATION NUMBER: US 09/390,039
; PRIOR FILING DATE: 1999-09-03
; PRIOR APPLICATION NUMBER: US 09/146,416
; PRIOR FILING DATE: 1998-09-03
; PRIOR APPLICATION NUMBER: US 10/024,036
; PRIOR FILING DATE: 2001-12-17
; PRIOR APPLICATION NUMBER: US 60/258,222
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: US 10/103,458
; PRIOR FILING DATE: 2002-03-22
; Remaining Prior Application data removed - See File Wrapper or PALM.
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 16:59:57 ; Search time 196.864 Seconds
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65.779 Million cell updates/sec

Title: US-10-754-446-7

Perfect score: 16

Sequence: 1 agaccaggccacat 16

Scoring table: IDENTITY NUC

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Total number of hits satisfying chosen parameters: 12077628

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Post-processing: Minimum Match 0%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	15	93.8	22	6 US-10-310-914A-1360219	Sequence 1360219,
4	15	93.8	25	6 US-10-310-914A-1360184	Sequence 1360184,
5	15	93.8	403278	6 US-10-995-561-13421	Sequence 13421, A
6	14.4	90.0	19	8 US-11-101-244-97619	Sequence 97619, A
7	14.4	90.0	19	8 US-11-101-244-97633	Sequence 97633, A
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9	14.4	90.0	19	9 US-11-083-784-97633	Sequence 97633, A
10	14.4	90.0	791	6 US-10-750-185-62019	Sequence 62019, A
11	14.4	90.0	791	6 US-10-750-623-62019	Sequence 62019, A
12	14.4	90.0	1214	7 US-11-112-908-134	Sequence 134, App
13	14.4	90.0	1889	6 US-10-821-234-37	Sequence 37, Appl
14	14.4	90.0	2215	7 US-11-112-908-144	Sequence 144, App
15	14.4	90.0	2453	6 US-10-750-185-36539	Sequence 36539, A
16	14.4	90.0	2453	6 US-10-750-623-36539	Sequence 36539, A
17	14.4	90.0	44848	7 US-11-106-672A-42	Sequence 42, Appl
18	14.4	90.0	54767	6 US-10-995-561-13357	Sequence 13357, A
19	14.4	90.0	158410	7 US-11-121-086-46	Sequence 46, Appl
20	14.4	90.0	164810	7 US-11-121-086-46	Sequence 4, Appl
21	14.4	90.0	180862	7 US-11-112-908-40	Sequence 40, Appl
22	14.4	90.0	189539	7 US-11-121-086-16	Sequence 16, Appl
23	14	87.5	201	7 US-11-124-368A-4143	Sequence 4143, Ap

24	14	87.5	496	6	US-10-750-185-37291	Sequence 37291, A
25	14	87.5	496	6	US-10-750-623-37291	Sequence 37291, A
26	14	87.5	600	7	US-11-136-527-6328	Sequence 6328, Ap
c 27	14	87.5	1038	6	US-10-512-109-16	Sequence 16, Appl
c 28	14	87.5	1223	6	US-10-512-109-3	Sequence 3, Appl
c 29	14	87.5	1223	6	US-10-512-109-14	Sequence 14, Appl
c 30	14	87.5	1816	6	US-10-750-185-29826	Sequence 29826, A
c 31	14	87.5	1816	6	US-10-750-623-29826	Sequence 29826, A
c 32	14	87.5	2378	6	US-10-821-234-117	Sequence 117, App
c 33	14	87.5	3884	6	US-10-131-826A-145	Sequence 145, App
c 34	14	87.5	10705	7	US-11-136-527-2232	Sequence 2232, Ap
c 35	14	87.5	20945	6	US-10-995-561-13463	Sequence 13463, A
c 36	14	87.5	23082	6	US-10-995-561-13457	Sequence 13457, A
c 37	14	87.5	27615	7	US-11-136-527-514	Sequence 514, App
c 38	14	87.5	94510	6	US-10-995-561-13332	Sequence 13332, A
c 39	14	87.5	103660	6	US-10-995-561-13253	Sequence 13253, A
c 40	14	87.5	197096	7	US-11-121-086-107	Sequence 107, App
c 41	13.4	83.8	19	8	US-10-310-914A-1165996	Sequence 1165996,
c 42	13.4	83.8	19	8	US-11-101-244-798463	Sequence 798463,
c 43	13.4	83.8	19	9	US-11-083-784-798463	Sequence 798463,
c 44	13.4	83.8	21	6	US-10-310-914A-1130386	Sequence 1130386,
c 45	13.4	83.8	23	6	US-10-310-914A-223914	Sequence 223914,

ALIGNMENTS

RESULT 1

US-11-112-908-22
; Sequence 22, Application US/11112908
; Publication No. US20050260659A1
; GENERAL INFORMATION:
; APPLICANT: Harris, Cole
; TITLE OF INVENTION: Breast Cancer Biomarkers
; FILE REFERENCE: 04-164-US
; CURRENT APPLICATION NUMBER: US/11/112,908
; CURRENT FILING DATE: 2005-04-22
; PRIOR APPLICATION NUMBER: US 60/564,758
; PRIOR FILING DATE: 2004-04-23
; PRIOR APPLICATION NUMBER: US 60/575,978
; PRIOR FILING DATE: 2004-06-01
; PRIOR APPLICATION NUMBER: US 60/631,702
; PRIOR FILING DATE: 2004-11-30
; PRIOR APPLICATION NUMBER: US 60/633,826
; PRIOR FILING DATE: 2004-12-07
; NUMBER OF SEQ ID NOS: 511
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 22
; LENGTH: 172147
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-112-908-22

Query Match 100.0%; Score 16; DB 7; Length 172147;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACCCAGGCCACAT 16

Db 116659 AGACCCAGGCCACAT 116674

RESULT 2

US-11-112-908-23
; Sequence 23, Application US/11112908
; Publication No. US20050260659A1
; GENERAL INFORMATION:
; APPLICANT: Harris, Cole
; TITLE OF INVENTION: Breast Cancer Biomarkers
; FILE REFERENCE: 04-164-US
; CURRENT APPLICATION NUMBER: US/11/112,908

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; CURRENT FILING DATE: 2005-04-22
; PRIOR APPLICATION NUMBER: US 60/564,758
; PRIOR FILING DATE: 2004-04-23
; PRIOR APPLICATION NUMBER: US 60/575,978
; PRIOR FILING DATE: 2004-06-01
; PRIOR APPLICATION NUMBER: US 60/631,702
; PRIOR FILING DATE: 2004-11-30
; PRIOR APPLICATION NUMBER: US 60/633,826
; PRIOR FILING DATE: 2004-12-07
; NUMBER OF SEQ ID NOS: 511
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 23
; LENGTH: 188682
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-112-908-23

Query Match      100.0%; Score 16; DB 7; Length 188682;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGACCCAGGCCCCACAT 16
Db      66280 AGACCCAGGCCCCACAT 66295

RESULT 3
US-10-310-914A-1360219
; Sequence 1360219, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvazat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 1360219
; LENGTH: 22
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-1360219

Query Match      93.8%; Score 15; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 2.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGACCCAGGCCCCACA 15
Db      8 AGACCCAGGCCCCACA 22

RESULT 4
US-10-310-914A-1360184
; Sequence 1360184, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvazat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 1360184
; LENGTH: 25
; TYPE: RNA
; ORGANISM: Human
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US-10-310-914A-1360184

Query Match      93.8%; Score 15; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 2e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGACCCAGGCCCCACA 15
Db      2 AGACCCAGGCCCCACA 16

RESULT 5
US-10-995-561-13421
; Sequence 13421, Application US/10995561
; Publication No. US20050272054A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; TITLE OF INVENTION: DETECTION AND USES THEREOF
; FILE REFERENCE: CL001559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13421
; LENGTH: 403278
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(403278)
; OTHER INFORMATION: n = A,T,C or G, or insertion/deletion polymorphism (see Tables 1-
US-10-995-561-13421

Query Match      93.8%; Score 15; DB 6; Length 403278;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGACCCAGGCCCCACA 15
Db      197280 AGACCCAGGCCCCACA 197294

RESULT 6
US-11-101-244-97619
; Sequence 97619, Application US/11101244
; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmason, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 13499US
; CURRENT APPLICATION NUMBER: US/11/101,244
; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
; PRIOR FILING DATE: 2002-11-14
; NUMBER OF SEQ ID NOS: 1591911
; SOFTWARE: Proprietary
; SEQ ID NO 97619
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Homo sapiens
US-11-101-244-97619

Query Match      90.0%; Score 14.4; DB 8; Length 19;
Best Local Similarity 87.5%; Pred. No. 4e+02;
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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488.341 Million cell updates/sec

Title: US-10-754-446-7

Perfect score: 16

Sequence: 1 agaccaggccacat 16

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Total number of hits satisfying chosen parameters: 2606114

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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8: /cgm2_6/ptodata/1/ina/RE COMB.seq.*
9: /cgm2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	16	100.0	2051	3	US-09-949-016-1712
3	16	100.0	15353	3	US-09-949-016-13454
4	15	93.8	2004	3	US-10-104-047-1222
5	15	93.8	56963	3	US-09-949-016-12966
6	15	93.8	56968	3	US-09-949-016-11888
7	15	93.8	254366	3	US-09-822-871-3
8	14.4	90.0	256	3	US-09-513-999C-25589
9	14.4	90.0	533	3	US-09-270-767-9339
10	14.4	90.0	533	3	US-09-270-767-24621
11	14.4	90.0	601	3	US-09-949-016-31585
12	14.4	90.0	601	3	US-09-949-016-31586
13	14.4	90.0	601	3	US-09-949-016-91141
14	14.4	90.0	601	3	US-09-949-016-94698
15	14.4	90.0	601	3	US-09-949-016-94699
16	14.4	90.0	601	3	US-09-949-016-151080
17	14.4	90.0	601	3	US-09-949-016-151081
18	14.4	90.0	1224	3	US-09-581-105-7
19	14.4	90.0	2502	2	US-08-073-384C-7
20	14.4	90.0	2502	2	US-08-254-359A-7
21	14.4	90.0	2502	2	US-08-483-043-7
22	14.4	90.0	2502	2	US-08-481-238-7
23	14.4	90.0	2502	2	US-08-471-066B-7
24	14.4	90.0	2502	2	US-08-484-956-7

25	14.4	90.0	2502	2	US-08-757-653-7	Sequence 7, Appli
26	14.4	90.0	2502	2	US-08-599-491-7	Sequence 7, Appli
27	14.4	90.0	2502	2	US-08-756-386-7	Sequence 7, Appli
28	14.4	90.0	2502	2	US-08-823-516-7	Sequence 7, Appli
29	14.4	90.0	2502	3	US-08-682-853A-7	Sequence 7, Appli
30	14.4	90.0	2502	3	US-08-759-038-7	Sequence 7, Appli
31	14.4	90.0	2502	3	US-08-758-314-7	Sequence 7, Appli
32	14.4	90.0	2502	3	US-09-350-309-7	Sequence 7, Appli
33	14.4	90.0	2502	3	US-08-520-946-7	Sequence 7, Appli
34	14.4	90.0	2502	3	US-09-684-938-7	Sequence 7, Appli
35	14.4	90.0	2502	3	US-09-308-825A-7	Sequence 7, Appli
36	14.4	90.0	2502	3	US-09-758-282B-7	Sequence 7, Appli
37	14.4	90.0	2502	3	US-09-655-378A-7	Sequence 7, Appli
38	14.4	90.0	2502	3	US-09-940-244-7	Sequence 7, Appli
39	14.4	90.0	2502	3	US-09-333-145-7	Sequence 7, Appli
40	14.4	90.0	2502	3	US-09-577-304A-7	Sequence 7, Appli
41	14.4	90.0	2502	3	US-09-381-212-7	Sequence 7, Appli
42	14.4	90.0	2502	3	US-10-081-806-7	Sequence 7, Appli
43	14.4	90.0	2502	3	US-09-713-601A-7	Sequence 7, Appli
44	14.4	90.0	2601	3	US-10-104-047-1515	Sequence 1515, Ap
45	14.4	90.0	3496	3	US-08-660-451A-5	Sequence 5, Appli

ALIGNMENTS

RESULT 1

US-09-949-016-59085
; Sequence 59085, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 59085
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-59085

Query Match 100.0%; Score 16; DB 3; Length 601;
Best Local Similarity 100.0%; Pred. No. 1e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGACCCAGGCCACAT 16
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Db 449 AGACCCAGGCCACAT 464

RESULT 2

US-09-949-016-1712
; Sequence 1712, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20

; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1712
; LENGTH: 2051
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-1712

Query Match 100.0%; Score 16; DB 3; Length 2051;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
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Db 952 AGACCCAGGCCACAT 967

RESULT 3

US-09-949-016-13454
; Sequence 13454, Application US/09949016
; Patent No. 6812339

; GENERAL INFORMATION:

; APPLICANT: VENTER, J. Craig et al.

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

; CURRENT APPLICATION NUMBER: US/09/949,016

; PRIOR FILING DATE: 2000-04-14

; PRIOR APPLICATION NUMBER: 60/241,755

; PRIOR FILING DATE: 2000-10-20

; PRIOR APPLICATION NUMBER: 60/237,768

; PRIOR FILING DATE: 2000-10-03

; PRIOR APPLICATION NUMBER: 60/231,498

; PRIOR FILING DATE: 2000-09-08

; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13454
; LENGTH: 15353
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13454

Query Match 100.0%; Score 16; DB 3; Length 15353;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 16
|||
Db 7582 AGACCCAGGCCACAT 7597

RESULT 4

US-10-104-047-1222
; Sequence 1222, Application US/10104047
; Patent No. 6943241

; GENERAL INFORMATION:

; APPLICANT: HELIX RESEARCH INSTITUTE

; TITLE OF INVENTION: 6943241el full length cdNA

; FILE REFERENCE: H1-A0105

; CURRENT APPLICATION NUMBER: US/10/104,047

; CURRENT FILING DATE: 2002-03-25

; PRIOR APPLICATION NUMBER:

; PRIOR FILING DATE:

; NUMBER OF SEQ ID NOS: 4096

; SOFTWARE: PatentIn Ver. 2.1

; SEQ ID NO 1222

; LENGTH: 2004

; TYPE: DNA

; ORGANISM: Homo sapiens

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Best Local Similarity 100.0%; Pred. No. 4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

US-10-104-047-1222

Query Match 93.8%; Score 15; DB 3; Length 2004;
Best Local Similarity 100.0%; Pred. No. 3.4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 5

US-09-949-016-12966/c
; Sequence 12966, Application US/09949016
; Patent No. 6812339

; GENERAL INFORMATION:

; APPLICANT: VENTER, J. Craig et al.

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

; CURRENT APPLICATION NUMBER: US/09/949,016

; PRIOR FILING DATE: 2000-04-14

; PRIOR APPLICATION NUMBER: 60/241,755

; PRIOR FILING DATE: 2000-10-20

; PRIOR APPLICATION NUMBER: 60/237,768

; PRIOR FILING DATE: 2000-10-03

; PRIOR APPLICATION NUMBER: 60/231,498

; PRIOR FILING DATE: 2000-09-08

; NUMBER OF SEQ ID NOS: 207012

; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 12966

; LENGTH: 56963

; TYPE: DNA

; ORGANISM: Human

US-09-949-016-12966

Query Match 93.8%; Score 15; DB 3; Length 56963;
Best Local Similarity 100.0%; Pred. No. 4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGACCCAGGCCACAT 15
|||
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RESULT 6

US-09-949-016-11888/c
; Sequence 11888, Application US/09949016
; Patent No. 6812339

; GENERAL INFORMATION:

; APPLICANT: VENTER, J. Craig et al.

; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED

; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF

; CURRENT APPLICATION NUMBER: US/09/949,016

; PRIOR FILING DATE: 2000-04-14

; PRIOR APPLICATION NUMBER: 60/241,755

; PRIOR FILING DATE: 2000-10-20

; PRIOR APPLICATION NUMBER: 60/237,768

; PRIOR FILING DATE: 2000-10-03

; PRIOR APPLICATION NUMBER: 60/231,498

; PRIOR FILING DATE: 2000-09-08

; NUMBER OF SEQ ID NOS: 207012

; SOFTWARE: FastSeq for Windows Version 4.0

; SEQ ID NO 11888

; LENGTH: 56968

; TYPE: DNA

; ORGANISM: Human

US-09-949-016-11888

Query Match 93.8%; Score 15; DB 3; Length 56968;
Best Local Similarity 100.0%; Pred. No. 4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 785.792 Seconds
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1157.425 Million cell updates/sec

Title: US-10-754-446-5

Perfect score: 16

Sequence: 1 tctgcccacagtacct 16

Scoring table: IDENTITY_NUC

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Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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C 20	15	93.8	670	10	BV025162	BV025162 S212P6124
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C 22	15	93.8	4992	5	AJ720888	AJ720888 Gallus ga
C 23	15	93.8	5210	9	AK122332	AK122332 Mus muscu
C 24	15	93.8	6331	13	EMVRPVP	J04374 Eggplant mo
C 25	15	93.8	62000	8	AP001984	AP001984 Homo sapi
C 26	15	93.8	110000	1	CF000082	Continuation (25 o
C 27	15	93.8	110000	14	AC118875	Continuation (3 of
C 28	15	93.8	118103	14	AC009817	AC009817 Homo sapi
C 29	15	93.8	134697	8	AC114738	AC114738 Homo sapi
C 30	15	93.8	138317	8	AP003481	AP003481 Homo sapi
C 31	15	93.8	139512	8	HSJ1169J3	AL049652 Human DNA
C 32	15	93.8	140968	9	BX537299	BX537299 Mouse DNA
C 33	15	93.8	151698	14	AC132644	AC132644 Rattus no
C 34	15	93.8	152816	14	AC079739	AC079739 Homo sapi
C 35	15	93.8	156808	14	AC012407	AC012407 Homo sapi
C 36	15	93.8	161100	14	AC025659	AC025659 Homo sapi
C 37	15	93.8	161179	14	AC083954	AC083954 Homo sapi
C 38	15	93.8	161362	14	AL355503	AL355503 Homo sapi
C 39	15	93.8	167322	14	AC016788	AC016788 Homo sapi
C 40	15	93.8	168789	9	AC127318	AC127318 Mus muscu
C 41	15	93.8	171141	9	AC157651	AC157651 Mus muscu
C 42	15	93.8	172160	9	AC119996	AC119996 Mus muscu
C 43	15	93.8	172772	9	AC121923	AC121923 Mus muscu
C 44	15	93.8	173805	9	AC124548	AC124548 Mus muscu
C 45	15	93.8	175048	14	AC161042	AC161042 Mus muscu

ALIGNMENTS

RESULT 1
AF305572S3
LOCUS Homo sapiens mucolipin 1 (MCOLN1) gene, exons 3 and 4.
DEFINITION Homo sapiens mucolipin 1 (MCOLN1) gene, exons 3 and 4.
ACCESSION AF305574
VERSION AF305574.1 GI:11991200
KEYWORDS
SEGMENT
SOURCE
3 of 8
Homo sapiens (human)

REFERENCE
1 (bases 1 to 790)
Bargal,R., Avidan,N., Ben-Asher,E., Olender,Z., Zeigler,M.,
Frumkin,A., Raas-Rothschild,A., Glusman,G., Lancet,D. and Bach,G.
Identification of the gene causing mucopolidiosis type IV
Nat. Genet. 26 (1), 118-123 (2000)
REFERENCE
2 (bases 1 to 790)
Bargal,R., Avidan,N., Ben-Asher,E., Olender,A., Zeigler,M.,
Frumkin,A., Raas-Rothschild,A., Glusman,G., Lancet,D. and Bach,G.
Direct Submission
Submitted (13-SEP-2000) Molecular Genetics, The Weizmann Institute
of Science, P. O. Box 26, Rehovot 76100, Israel
FEATURES
Location/Qualifiers
1..790
/organism="Homo sapiens"
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/db_xref="taxon:9606"
267..434
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/number=3
589..757
/gene="MCOLN1"
/number=4

ORIGIN

Query Match 100.0%; Score 16; DB 8; Length 790;
Best Local Similarity 100.0%; Pred. No. 1.6e+02; Indels 0; Gaps 0;
Matches 16; Conservative 0; Mismatches 0;

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 1789.31 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-5
Perfect score: 16
Sequence: 1 tctgccacagctacct 16

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : EST.*
1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_hc.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_est7.*
9: gb_gss1.*
10: gb_gss2.*
11: gb_gss3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	16	100.0	154	9 BH271054	BH271054 CH230-162
2	16	100.0	480	2 BB856429	BB856429 BB856429
3	16	100.0	525	9 AQ669413	AQ669413 HS_5407_B
C 4	16	100.0	696	1 AJ727930	AJ727930 AJ727930
5	16	100.0	745	1 AJ727936	AJ727936 AJ727936
6	16	100.0	762	5 BU306972	BU306972 603612510
7	16	100.0	774	8 DR431852	DR431852 nax48f12
8	16	100.0	806	5 BU383135	BU383135 603961896
9	16	100.0	1008	5 BU420692	BU420692 603231561
C 10	15	93.8	224	8 CV972084	CV972084 LRRGE0208
C 11	15	93.8	269	8 CV972398	CV972398 LRRGE0239
12	15	93.8	279	1 BB007743	BB007743 BB007743
13	15	93.8	279	8 CV972703	CV972703 LRRGE0270
14	15	93.8	281	8 CV888115	CV888115 LRRGE0000
C 15	15	93.8	361	8 CV972086	CV972086 LRRGE0208
16	15	93.8	388	5 BY619013	BY619013 BY619013
C 17	15	93.8	420	9 BH273644	BH273644 CH230-70E
C 18	15	93.8	436	8 CV888113	CV888113 LRRGE0000
19	15	93.8	475	8 CV888114	CV888114 LRRGE0000
C 20	15	93.8	492	9 AZ822326	AZ822326 2M0095J01
C 21	15	93.8	501	8 BH117609	BH117609 RPCI-24-2
22	15	93.8	507	9 AZ768935	AZ768935 1M0569J09

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23 15 93.8 517 9 AZ585292
24 15 93.8 525 9 AQ357179
c 25 15 93.8 591 10 CE537462
26 15 93.8 595 2 BG404774
c 27 15 93.8 599 1 AI412276
28 15 93.8 625 7 CK620962
29 15 93.8 672 10 CE809412
30 15 93.8 714 5 BY764892
31 15 93.8 727 5 BU724937
c 32 15 93.8 727 5 BU139411
c 33 15 93.8 732 1 AJ724931
c 34 15 93.8 734 10 CE812263
c 35 15 93.8 745 11 CR081738
c 36 15 93.8 753 10 AG457089
c 37 15 93.8 763 11 CR228261
c 38 15 93.8 784 3 BI331708
c 39 15 93.8 784 7 CF924020
c 40 15 93.8 790 7 CN234992
c 41 15 93.8 797 7 CK315394
c 42 15 93.8 870 11 CR118310
c 43 15 93.8 999 2 BF976532
c 44 15 93.8 1001 5 BU192451
c 45 15 93.8 1184 7 CK167560

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ALIGNMENTS

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RESULT 1
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LOCUS BH271054
DEFINITION CH230-162L17.TJ CHORI-230 Segment 1 Rattus norvegicus genomic clone
CH230-162L17, genomic survey sequence.
ACCESSION BH271054
VERSION BH271054.1 GI:17183456
KEYWORDS GSS
SOURCE Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridea; Muridae; Murinae; Rattus.
1 (bases 1 to 154)
Zhao,S., Shetty,J., Shatsman,S., Tsengay,G., Geer,K.,
Shvartsbeyn,A., Gebregeorgis,E., Overton,L., Russeil,D., Chen,D.,
Riggs,F., de Jong,P. and Fraser,C.M.
Rat BAC End Sequences from Library CHORI-230 EcoRI segment
Unpublished (1999)
Other_GSSs: CH230-162L17.TV
Contact: Shaying Zhao
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: szhao@tigr.org
Clones are derived from the rat BAC library CHORI-230
(http://www.chori.org/bacpac/rac230.htm). For BAC library
availability, please contact Pieter de Jong (pdejong@mail.cho.org).
Clones may be purchased from BACPAC Resources
(http://www.chori.org/bacpac/or ering_information.htm). BAC end
page: http://www.tigr.org/tadb/bac_ends/rat/bac_end_intro.html
Plate: 162 row: L column: 17
Seq primer: SP6
Class: BAC ends.
Location/Qualifiers
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/sex="Female"
/cell_type="Brain"

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/clone_lib="CHORI-230 Segment 1"
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 Pieter de Jong"

ORIGIN

Query Match 100.0%; Score 16; DB 9; Length 154;
 Best Local Similarity 100.0%; Pred. No. 1.4e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCTGCCCACAGTACCT 16
 |||||
 Db 150 TCTGCCCACAGTACCT 135

RESULT 2

BB856429

LOCUS BB856429 RIKEN full-length enriched, B16 F10Y cells linear EST 26-NOV-2001
 DEFINITION cDNA clone G370034110 5', mRNA sequence.

ACCESSION BB856429

VERSION BB856429.1 GI:17097883

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM

Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
 Sciurognathi; Muridae; Muridae; Murinae; Mus.

REFERENCE

AUTHORS

1 (bases 1 to 480)
 Akimura,T., Arakawa,T., Carninci,P., Furuno,M., Hanagaki,T.,
 Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane,T., Imotani,K.,
 Ishii,Y., Ito,M., Kawai,J., Kojima,Y., Konno,H., Kouda,M.,
 Matsuyama,T., Nakamura,M., Nishi,K., Nomura,K., Numasaki,R.,
 Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sakazume,N.,
 Sasaki,D., Sato,K., Shibata,K., Shinagawa,A., Shiraki,T.,
 Sogabe,Y., Suzuki,H., Tagawa,A., Takahashi,F., Takaku-Akaihi,S.,
 Tanaka,T., Tomaru,A., Tota,T., Watahiki,A., Yasunishi,A.,
 Muramatsu,M. and Hayashizaki,Y.

RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura,T., et al.
 2001)

TITLE

JOURNAL

COMMENT

Unpublished (2001)
 Contact: Yoshihide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center(GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suenho-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216

Email: genome-resgsc.riken.jp, URL:http://genome.gsc.riken.jp/
 Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K.,
 Itoh,M., Konno,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new
 genes. Genome Res. 10 (10), 1617-1630 (2000)
 wagi,K., Fujiwara,S., Inoue,K., Togawa,Y., Izawa,M., Ohara,E.,
 Watahiki,M., Yoneda,Y., Ishikawa,T., Ozawa,K., Tanaka,T.,
 Matsuura,S., Kawai,J., Okazaki,Y., Muramatsu,M., Inoue,Y., Kira,A.
 and Hayashizaki,Y.

RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multipipillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Konno,H., Fukunishi,Y., Shibata,K., Itoh,M., Carninci,P.,
 Sugahara,Y. and Hayashizaki,Y.

Computer-based methods for the mouse full-length cDNA
 encyclopedia: real-time sequence clustering for construction of a
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
 Please visit our web site (http://genome.gsc.riken.go.jp) for
 further details.

e mouse tissues.

FEATURES

source

1. .480
 /organism="Mus musculus"
 /mol_type="mRNA"

/strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="G370034110"
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ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 1.6e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCTGCCCACAGTACCT 16
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 Db 366 TCTGCCCACAGTACCT 381

RESULT 3

AQ669413

LOCUS

DEFINITION AQ669413 525 bp DNA linear GSS 24-JUN-1999
 HS 5407_B1_F10_T7A RPCI-11 Human Male BAC Library Homo sapiens
 genomic clone Plate=983 Col=19 Row=L, genomic survey sequence.

ACCESSION AQ669413

VERSION AQ669413.1 GI:5202159

KEYWORDS GSS.

SOURCE Homo sapiens (human)

ORGANISM

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
 Homnidae; Homo.

REFERENCE

AUTHORS

1 (bases 1 to 525)
 Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
 Hood,L.

Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome

JOURNAL

PUBMED

COMMENT

10449764
 Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887

Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieteredejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
 or from Research Genetics (info@resgen.com). BAC end Web Server:
 http://www.htsc.washington.edu

Plate: 983 row: L column: 19
 Seq primer: T7
 Class: BAC ends

High quality sequence stop: 525.

Location/Qualifiers

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/mol_type="genomic DNA"

/db_xref="taxon:9606"

/clone="Plate=983 Col=19 Row=L"

/sex="male"

/clone_lib="RPCI-11 Human Male BAC Library"

/note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI;
 Male blood DNA was isolated from one randomly chosen donor
 and partially digested with a combination of EcoRI and
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FEATURES

source

1. .525
 /organism="Homo sapiens"
 /mol_type="genomic DNA"

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/note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI;
 Male blood DNA was isolated from one randomly chosen donor
 and partially digested with a combination of EcoRI and
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 pBACe3.6 vector at EcoRI sites"

ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 1.6e+03;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 217.088 Seconds
(without alignments)
491.207 Million cell updates/sec

Title: US-10-754-446-5

Perfect score: 16

Sequence: 1 tctgcccacagtaacct 16

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 3332346308 residues

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: Geneseqn2000s.*
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5: Geneseqn2001bs.*
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11: Geneseqn2003ds.*
12: Geneseqn2004as.*
13: Geneseqn2004bs.*
14: Geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	16	100.0	16	14 AEB28961	Aeb28961 Human MCO
2	16	100.0	11365	4 AAK73827	Aak73827 Human imm
3	16	100.0	13270	9 ADB84283	Adb84283 Human muc
4	16	100.0	20046	4 AAK73826	Aak73826 Human imm
5	15	93.8	177531	8 ACF62732	Acf62732 Cancer ba
6	15	93.8	177531	8 ADB20847	Adb20847 MRPI base
7	15	93.8	177531	10 ADB87936	Adb87936 Human UGT
8	15	93.8	177531	10 ADB96919	Adb96919 Human MDR
9	15	93.8	177531	10 ADB92110	Adb92110 Human MDR
10	15	93.8	177531	10 ADH74617	Adh74617 Human BAC
11	14.4	90.0	405	4 AAL35222	Aal35222 Human mus
12	14.4	90.0	405	8 ABX58210	Abx58210 cDNA enco
13	14.4	90.0	405	12 ADJ27937	Adj27937 Human mus
c 14	14.4	90.0	465	9 ACH32754	Ach32754 Human end
c 15	14.4	90.0	501	10 ADB49981	Adb49981 Primary r
c 16	14.4	90.0	548	13 ADQ50585	Adq50585 Novel can
c 17	14.4	90.0	570	4 AAL13764	Aal13764 Human bre
c 18	14.4	90.0	789	2 AAV65231	Aav65231 DNA seque
c 19	14.4	90.0	835	6 ABU90105	Abu90105 Human pol

c	20	14.4	90.0	897	11	ACN83833	Acn83833 Breast ca
	21	14.4	90.0	974	6	AAD31825	Aad31825 Human pan
	22	14.4	90.0	1044	13	ADS56881	Ads56881 Bacterial
c	23	14.4	90.0	1692	4	AAK79833	Aak79833 Human imm
	24	14.4	90.0	1849	5	AAS83682	Aas83682 DNA enco
	25	14.4	90.0	2214	13	ADU01699	Adu01699 Novel hum
	26	14.4	90.0	2228	6	ABQ99407	Abq99407 Human cod
	27	14.4	90.0	2297	12	ADQ55977	Adq55977 T cell ac
	28	14.4	90.0	2317	6	ABS76414	Abs76414 cDNA enco
	29	14.4	90.0	2317	6	ABV99386	Abv99386 Human NOV
	30	14.4	90.0	2317	13	ADU06214	Adu06214 Novel bro
	31	14.4	90.0	2380	4	AAL36812	Aal36812 Human mus
	32	14.4	90.0	2380	8	ABX59800	Abx59800 cDNA enco
	33	14.4	90.0	2380	12	ADJ30550	Adj30550 Human mus
	34	14.4	90.0	2394	4	AAL36811	Aal36811 Human mus
	35	14.4	90.0	2394	8	ABX59799	Abx59799 cDNA enco
	36	14.4	90.0	2394	12	ADJ30549	Adj30549 Human mus
	37	14.4	90.0	2452	6	AAI71445	Aai71445 TNFR/NGFR
	38	14.4	90.0	2569	10	ADB62298	Adb62298 Human cDN
	39	14.4	90.0	2788	4	AAH17843	Aah17843 Human cDN
	40	14.4	90.0	2816	13	ACN38987	Acn38987 Tumour-as
	41	14.4	90.0	3025	2	AAQ13337	Aaq13337 D1 dopami
	42	14.4	90.0	3025	2	AAT63657	Aat63657 D1 dopami
c	43	14.4	90.0	3778	6	ABZ11278	Abz11278 Human pol
c	44	14.4	90.0	3778	12	ADM43796	Adm43796 Novel hum
c	45	14.4	90.0	4376	10	ADF50290	Adf50290 Human PFM

ALIGNMENTS

RESULT 1

AEB28961
ID AEB28961 standard; DNA; 16 BP.
XX
AC AEB28961;
XX
DT 22-SEP-2005 (first entry)
XX
DE Human MCOLN1 gene probe IVS WT SEQ ID NO:5.
XX
KW mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; probe;
KW ss.
XX
OS Homo sapiens.
PN
PN US2005153300-A1.
PD 14-JUL-2005.
XX
PF 09-JAN-2004; 2004US-00754446.
XX
PR 09-JAN-2004; 2004US-00754446.
XX
XX (QUES-) QUEST DIAGNOSTICS INC.
XX
XX Sun W, Hantash F;
XX
XX WPI; 2005-521160/53.

Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by amplifying the nucleic acid, detecting amplified product with labeled oligonucleotide probes via a change in fluorescence which indicates the presence of an ML IV mutant.

Claim 3; SEQ ID NO 5; 15pp; English.

The invention relates to a method (M1) for detecting the presence of mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1) involves contacting the nucleic acid with oligonucleotide primers and probes, conducting amplification by temperature cycling and monitoring the accumulation of amplified nucleic acid by detecting an increase in donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucolin-1
 CC (MCOLN1) gene; and (2) a kit (KI) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a probe for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

XX
 SQ Sequence 16 BP; 3 A; 7 C; 2 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 16; DB 14; Length 16;

Best Local Similarity 100.0%; Pred. No. 1.3e+02; Mismatches 0; Indels 0; Gaps 0;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCTGCCACAGTACCT 16

Db 1 TCTGCCACAGTACCT 16

RESULT 2

AAK73827

ID AAK73827 standard; DNA; 11365 BP.

XX AC AAK73827;

XX DT 07-NOV-2001 (first entry)

XX DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:28639.

XX KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;

XX KW Cytostatic; Gene therapy; vaccine; metastasis; ds.

XX OS Homo sapiens.

XX PN WO200157182-A2.

XX PD 09-AUG-2001.

XX PF 17-JAN-2001; 2001WO-US001354.

XX PR 31-JAN-2000; 2000US-0179065P.

PR 04-FEB-2000; 2000US-0180628P.

PR 24-FEB-2000; 2000US-0184664P.

PR 02-MAR-2000; 2000US-0186350P.

PR 16-MAR-2000; 2000US-0189874P.

PR 17-MAR-2000; 2000US-0190076P.

PR 16-APR-2000; 2000US-0198123P.

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Total number of hits satisfying chosen parameters: 19587084

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	16	100.0	653	4	US-09-925-065A-696500
4	16	100.0	13270	3	US-09-851-494B-1
5	15.6	97.5	600	9	US-10-972-079-69742
6	15.6	97.5	600	9	US-10-972-079-69743
7	15.6	97.5	600	9	US-10-972-079-69744
8	15.6	97.5	600	9	US-10-972-079-69745
9	15	93.8	310	7	US-10-242-535A-32082
10	15	93.8	310	7	US-10-085-783A-32082
11	15	93.8	533	5	US-10-027-632-224114
12	15	93.8	533	6	US-10-027-632-224114
13	15	93.8	177531	8	US-10-484-577-660
14	14.4	90.0	25	7	US-10-719-956-630855
15	14.4	90.0	275	9	US-10-756-149-4182
16	14.4	90.0	295	8	US-10-674-124A-10971
17	14.4	90.0	297	3	US-09-783-590-8724
18	14.4	90.0	312	6	US-10-062-674-400
19	14.4	90.0	388	3	US-09-783-590-8696
20	14.4	90.0	390	3	US-09-783-590-8711
21	14.4	90.0	390	3	US-09-783-590-8711
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25 14.4 90.0 405 6 US-10-242-515-564 Sequence 564, App
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27 14.4 90.0 452 4 US-09-925-065A-256794 Sequence 256794,
28 14.4 90.0 452 4 US-09-925-065A-256795 Sequence 256795,
29 14.4 90.0 452 4 US-09-925-065A-256796 Sequence 256796,
30 14.4 90.0 452 3 US-09-918-995-19966 Sequence 19966, A
31 14.4 90.0 533 4 US-09-925-065A-112177 Sequence 112177,
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33 14.4 90.0 563 4 US-09-925-065A-197601 Sequence 197601,
34 14.4 90.0 585 5 US-10-027-632-229737 Sequence 229737,
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ALIGNMENTS

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; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: SUN, WEIMIN
; APPLICANT: HANTASH, PERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; TITLE OF INVENTION: MUCOLIPIDOSIS IV MUTATIONS
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10/754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: Patent In Ver. 3.2
; SEQ ID NO 5
; LENGTH: 16
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Probe
US-10-754-446-5

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Best Local Similarity 100.0%; Pred. No. 90;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 TCTGCCCCACAGTACCT 16

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; Publication No. US20050228172A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single
; TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
; FILE REFERENCE: 198827.135
; CURRENT APPLICATION NUMBER: US/09/925,065A
; CURRENT FILING DATE: 2001-08-08
; PRIOR APPLICATION NUMBER: US 60/243,096
; PRIOR FILING DATE: 2000-10-24
; PRIOR APPLICATION NUMBER: US 60/252,147
; PRIOR FILING DATE: 2000-11-20

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; PRIOR APPLICATION NUMBER: US 60/250,092
; PRIOR FILING DATE: 2000-11-30
; PRIOR APPLICATION NUMBER: US 60/261,766
; PRIOR FILING DATE: 2001-01-16
; PRIOR APPLICATION NUMBER: US 60/289,846
; PRIOR FILING DATE: 2001-05-09
; NUMBER OF SEQ ID NOS: 957086
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 696499
; LENGTH: 653
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-925-065A-696499

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Best Local Similarity 100.0%; Pred. No. 73;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 137 TCTGCCCCACAGTACCT 152

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; Sequence 696500, Application US/09925065A
; Publication No. US20030228172A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single
; FILE REFERENCE: 108827.135
; CURRENT APPLICATION NUMBER: US/09/925,065A
; PRIOR FILING DATE: 2001-08-08
; PRIOR APPLICATION NUMBER: US 60/243,096
; PRIOR FILING DATE: 2000-10-24
; PRIOR APPLICATION NUMBER: US 60/252,147
; PRIOR FILING DATE: 2000-11-20
; PRIOR APPLICATION NUMBER: US 60/250,092
; PRIOR FILING DATE: 2000-11-30
; PRIOR APPLICATION NUMBER: US 60/261,766
; PRIOR FILING DATE: 2001-01-16
; PRIOR APPLICATION NUMBER: US 60/289,846
; PRIOR FILING DATE: 2001-05-09
; NUMBER OF SEQ ID NOS: 957086
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 696500
; LENGTH: 653
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-925-065A-696500

Query Match 100.0%; Score 16; DB 4; Length 653;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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US-09-851-494B-1
; Sequence 1, Application US/09851494B
; Publication No. US20030064363A1
; GENERAL INFORMATION:
; APPLICANT: ML4 Foundation
; APPLICANT: Goldin, Ehud
; APPLICANT: Slaugenhaupt, Susan A.
; APPLICANT: Sun, Mei
; APPLICANT: Acierno, James S.
; TITLE OF INVENTION: A Gene Encoding A New TRP Channel is Mutated in Mucopolipidosis IV
; FILE REFERENCE: 3394/1H57U51

; CURRENT APPLICATION NUMBER: US/09/851,494B
; CURRENT FILING DATE: 2002-07-12
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
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; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-851-494B-1

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Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 5525 TCTGCCCCACAGTACCT 5540

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US-10-972-079-69742
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; Publication No. US20050153317A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: ROSENFELD, David
; APPLICANT: KERR, Richard
; APPLICANT: BATES, Stephen
; APPLICANT: HOLM, Tom
; TITLE OF INVENTION: METHODS & SYSTEMS FOR INFERRING TRAITS TO BREED & MANAGE NON-BEEP
; FILE REFERENCE: MM1110-2
; CURRENT APPLICATION NUMBER: US/10/972,079
; CURRENT FILING DATE: 2004-10-22
; PRIOR APPLICATION NUMBER: US 60/514,333
; PRIOR FILING DATE: 2003-10-24
; NUMBER OF SEQ ID NOS: 96631
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 69742
; LENGTH: 600
; TYPE: DNA
; ORGANISM: Chicken 19866894342124_1
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US-10-972-079-69743
; Sequence 69743, Application US/10972079
; Publication No. US20050153317A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: ROSENFELD, David
; APPLICANT: KERR, Richard
; APPLICANT: BATES, Stephen
; APPLICANT: HOLM, Tom
; TITLE OF INVENTION: METHODS & SYSTEMS FOR INFERRING TRAITS TO BREED & MANAGE NON-BEEP
; FILE REFERENCE: MM1110-2
; CURRENT APPLICATION NUMBER: US/10/972,079
; CURRENT FILING DATE: 2004-10-22
; PRIOR APPLICATION NUMBER: US 60/514,333
; PRIOR FILING DATE: 2003-10-24
; NUMBER OF SEQ ID NOS: 96631

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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C 18	14	87.5	159138	6	US-10-995-561-13230	Sequence 13230, A	
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C 21	13.4	83.8	25	7	US-11-121-849-638454	Sequence 638454, A	
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; GENERAL INFORMATION:
; APPLICANT: Bentwach, Isaac
; APPLICANT: Shiler, Kuzat
; TITLE OF INVENTION: Bioinformatically deter
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 10301
; LENGTH: 69
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-10301

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Matches 11: Conservative 4; Mismatches 1; Indels

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Publication No. US2006003322A1
GENERAL INFORMATION:
APPLICANT: Bentwich, Isaac
APPLICANT: Shiler, Kuvzat
TITLE OF INVENTION: Bioinformatically deter
TITLE OF INVENTION: uses thereof
FILE REFERENCE: 06087, 0200.CPUS01
CURRENT APPLICATION NUMBER: US/10/310,914A
CURRENT FILING DATE: 2002-12-06
NUMBER OF SEQ ID NOS: 1388402
SOFTWARE: Patent In version 3.3
SEQ ID NO 16743
LENGTH: 69
TYPE: RNA

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Sequence 43019, A
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Sequence 32683, A
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Sequence 27624, A
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Sequence 40981, A

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; ORGANISM: Human
US-10-310-914A-16743

Query Match          90.0%; Score 14.4; DB 6; Length 69;
Best Local Similarity 93.8%; Pred. No. 73;
Matches 11; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

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Db 28 UCUGCACACAGUACCU 43

RESULT 3
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; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10750,185
; PRIOR FILING DATE: 2003-12-31
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 61765
; LENGTH: 1126
; TYPE: DNA
; ORGANISM: Bovine 19866881607649
US-10-750-185-61765

Query Match          90.0%; Score 14.4; DB 6; Length 1126;
Best Local Similarity 93.8%; Pred. No. 80;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
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Db 599 TCTGCCCCACAGTACCT 584

RESULT 4
US-10-750-623-61765/c
; Sequence 61765, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10750,623
; PRIOR FILING DATE: 2003-12-31
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 61765
; LENGTH: 1126
; TYPE: DNA
; ORGANISM: Bovine 19866881607649
US-10-750-623-61765

Query Match          90.0%; Score 14.4; DB 6; Length 1126;
Best Local Similarity 93.8%; Pred. No. 81;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
   ||||| ||||| |||||
Db 1253 TCTGCCCCACAGTACCT 1238

RESULT 5
US-10-750-185-48943/c
; Sequence 48943, Application US/10750185
; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10750,185
; PRIOR FILING DATE: 2003-12-31
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 48943
; LENGTH: 2095
; TYPE: DNA
; ORGANISM: Bovine 19866881228818
US-10-750-185-48943

Query Match          90.0%; Score 14.4; DB 6; Length 2095;
Best Local Similarity 93.8%; Pred. No. 81;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
   ||||| ||||| |||||
Db 1253 TCTGCCCCACAGTACCT 1238

RESULT 6
US-10-750-623-48943/c
; Sequence 48943, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10750,623
; PRIOR FILING DATE: 2003-12-31
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIN version 3.1
; SEQ ID NO 48943
; LENGTH: 2095
; TYPE: DNA
; ORGANISM: Bovine 19866881228818
US-10-750-623-48943

Query Match          90.0%; Score 14.4; DB 6; Length 2095;
Best Local Similarity 93.8%; Pred. No. 81;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
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GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 16:32:47 ; Search time 58.24 Seconds
(without alignments)
488.341 Million cell updates/sec

Title: US-10-754-446-5

Perfect score: 16

Sequence: 1 tctgcccacagtacct 16

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Gapop 10.0 , Gapext 1.0

Searched: 1303057 seqs, 888780828 residues

Total number of hits satisfying chosen parameters: 2606114

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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- 8: /cgn2_6/ptodata/1/ina/RE COMB.seq.*
- 9: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	14.4	90.0	601	3	US-09-949-016-22227 Sequence 22227, A
5	14.4	90.0	601	3	US-09-949-016-22228 Sequence 22228, A
6	14.4	90.0	601	3	US-09-949-016-22229 Sequence 22229, A
7	14.4	90.0	601	3	US-09-949-016-22230 Sequence 22230, A
8	14.4	90.0	601	3	US-09-949-016-22231 Sequence 22231, A
9	14.4	90.0	601	3	US-09-949-016-22232 Sequence 22232, A
10	14.4	90.0	601	3	US-09-949-016-22233 Sequence 22233, A
11	14.4	90.0	601	3	US-09-949-016-38209 Sequence 38209, A
12	14.4	90.0	601	3	US-09-949-016-38210 Sequence 38210, A
13	14.4	90.0	601	3	US-09-949-016-38211 Sequence 38211, A
14	14.4	90.0	601	3	US-09-949-016-51333 Sequence 51333, A
15	14.4	90.0	601	3	US-09-949-016-51334 Sequence 51334, A
16	14.4	90.0	601	3	US-09-949-016-51335 Sequence 51335, A
17	14.4	90.0	601	3	US-09-949-016-51336 Sequence 51336, A
18	14.4	90.0	601	3	US-09-949-016-51337 Sequence 51337, A
19	14.4	90.0	601	3	US-09-949-016-51338 Sequence 51338, A
20	14.4	90.0	601	3	US-09-949-016-51339 Sequence 51339, A
21	14.4	90.0	601	3	US-09-949-016-51340 Sequence 51340, A
22	14.4	90.0	601	3	US-09-949-016-68413 Sequence 68413, A
23	14.4	90.0	601	3	US-09-949-016-68414 Sequence 68414, A
24	14.4	90.0	601	3	US-09-949-016-68415 Sequence 68415, A

C 25	14.4	90.0	601	3	US-09-949-016-169541 Sequence 169541, A
C 26	14.4	90.0	601	3	US-09-949-016-169542 Sequence 169542, A
C 27	14.4	90.0	601	3	US-09-949-016-169543 Sequence 169543, A
C 28	14.4	90.0	789	3	US-08-986-765-4 Sequence 4, Appli
C 29	14.4	90.0	2569	3	US-10-104-047-452 Sequence 452, App
C 30	14.4	90.0	2883	3	US-09-614-231A-371 Sequence 371, Appli
C 31	14.4	90.0	3025	2	US-08-444-734A-1 Sequence 1, Appli
C 32	14.4	90.0	3778	3	US-09-799-451-160 Sequence 160, App
C 33	14.4	90.0	4376	3	US-10-200-012-15 Sequence 15, Appli
C 34	14.4	90.0	4521	3	US-09-533-494A-18 Sequence 18, Appli
C 35	14.4	90.0	8461	3	US-09-949-016-13428 Sequence 13428, A
C 36	14.4	90.0	12055	3	US-09-949-016-13233 Sequence 13233, A
C 37	14.4	90.0	67479	3	US-09-949-016-11804 Sequence 11804, A
C 38	14.4	90.0	71119	3	US-09-949-016-15358 Sequence 15358, A
C 39	14.4	90.0	77661	3	US-09-949-016-12770 Sequence 12770, A
C 40	14.4	90.0	77663	3	US-09-949-016-13751 Sequence 13751, A
C 41	14.4	90.0	121427	3	US-09-949-016-11950 Sequence 11950, A
C 42	14.4	90.0	121433	3	US-09-949-016-13230 Sequence 13230, A
C 43	14.4	90.0	160759	3	US-09-949-016-16514 Sequence 16514, A
C 44	14	87.5	601	3	US-09-949-016-117072 Sequence 117072, A
C 45	14	87.5	601	3	US-09-949-016-152388 Sequence 152388, A

ALIGNMENTS

RESULT 1

US-09-949-016-13454
; Sequence 13454, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13454
; LENGTH: 15353
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13454

Query Match 100.0%; Score 16; DB 3; Length 15353;
Best Local Similarity 100.0%; Pred. No. 44;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCTGCCACAGTACTT 16

DB 6125 TCTGCCACAGTACTT 6140

RESULT 2

US-09-949-016-13427/c
; Sequence 13427, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20

; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13427
; LENGTH: 110585
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(110585)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-13427

Query Match 100.0%; Score 16; DB 3; Length 110585;
Best Local Similarity 100.0%; Pred. No. 53;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
Db 7380 TCTGCCCCACAGTACCT 7365

RESULT 3

US-09-949-016-22226
; Sequence 22226 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22226
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-22226

Query Match 90.0%; Score 14.4; DB 3; Length 601;
Best Local Similarity 93.8%; Pred. No. 2.4e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
Db 77 TCTGCCCCACATACCT 92

RESULT 4

US-09-949-016-22227
; Sequence 22227 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768

; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22227
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-22227

Query Match 90.0%; Score 14.4; DB 3; Length 601;
Best Local Similarity 93.8%; Pred. No. 2.4e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
Db 94 TCTGCCCCACATACCT 109

RESULT 5

US-09-949-016-22228
; Sequence 22228 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22228
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-22228

Query Match 90.0%; Score 14.4; DB 3; Length 601;
Best Local Similarity 93.8%; Pred. No. 2.4e+02;
Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCTGCCCCACAGTACCT 16
Db 464 TCTGCCCCACATACCT 479

RESULT 6

US-09-949-016-22229
; Sequence 22229 Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0

GenCore version 5.1.6
Copyright (c) 1993 - 2006 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 1031.35 Seconds
(without alignments)
1157.425 Million cell updates/sec

Title: US-10-754-446-4

Perfect score: 21

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Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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C 3	21	100.0	155645	14 AC021153	AC021153 Homo sapi
C 4	21	100.0	171126	8 AC008878	AC008878 Homo sapi
C 5	18.4	87.6	1740	6 AX280021	AX280021 Sequence
C 6	18.4	87.6	1829	6 CQ719994	CQ719994 Sequence
C 7	18.4	87.6	2004	8 AF249319	AF249319 Homo sapi
C 8	18.4	87.6	2037	8 HSA293970	HSA293970 Homo sapi
C 9	18.4	87.6	2049	8 AK026102	AK026102 Homo sapi
C 10	18.4	87.6	2051	8 AF287269	AF287269 Homo sapi
C 11	18.4	87.6	2052	6 AX083508	AX083508 Sequence
C 12	18.4	87.6	2063	8 AX222673	AX222673 Homo sapi
C 13	18.4	87.6	2078	8 BC005149	BC005149 Homo sapi
C 14	18.4	87.6	2094	6 BD233734	BD233734 31 human
C 15	18.4	87.6	2095	6 AX280019	AX280019 Sequence
C 16	18.4	87.6	2272	8 HSA293659	HSA293659 Homo sapi
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C 18	18.4	87.6	195753	14 AC128091	AC128091 Rattus no

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C 20	18.4	87.6	224426	14 AC108247	AC108247 Rattus no
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C 22	18.4	87.6	241828	14 AC106575	AC106575 Rattus no
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C 26	18	85.7	110000	1 CP000050	Continuation (20 o
C 27	17.8	84.8	75511	14 AC153278	AC153278 Bos tauru
C 28	17.8	84.8	107113	14 AC142559	AC142559 Takifugu
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C 31	17.8	84.8	169812	14 AC150169	AC150169 Gallus ga
C 32	17.8	84.8	183916	8 AF111169	AF111169 Homo sapi
C 33	17.8	84.8	186330	8 AC007376	AC007376 Homo sapi
C 34	17.8	84.8	198278	14 AC150063	AC150063 Gallus ga
C 35	17.8	84.8	212150	14 AC130048	AC130048 Rattus no
C 36	17.8	84.8	224635	14 AC094848	AC094848 Rattus no
C 37	17.8	84.8	231625	14 AC107597	AC107597 Rattus no
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ALIGNMENTS

RESULT 1
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LOCUS AF30557285 579 bp DNA linear PRI 26-DEC-2000
DEFINITION Homo sapiens mucopolipin 1 (MCOLN1) gene, exons 6 and 7.
ACCESSION AF305576
VERSION AF305576.1 GI:11991202
KEYWORDS
SEGMENT
SOURCE 5 of 8
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1 (bases 1 to 579)
AUTHORS Bargal, R., Avidan, N., Ben-Asher, E., Olender, Z., Zeigler, M.,
Frumkin, A., Raas-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
TITLE Identification of the gene causing mucopolipidosis type IV
JOURNAL Nat. Genet. 26 (1), 118-123 (2000)
PUBMED 10973263
REFERENCE 2 (bases 1 to 579)
AUTHORS Bargal, R., Avidan, N., Ben-Asher, E., Olender, A., Zeigler, M.,
Frumkin, A., Raas-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
TITLE Direct Submission
JOURNAL Submitted (13-SEP-2000) Molecular Genetics, The Weizmann Institute
of Science, P. O. Box 26, Rehovot 76100, Israel
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exon
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Query Match 100.0%; Score 21; DB 8; Length 579;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      527 CTCACCGTGTGGAGACACT 507

RESULT 2
AF287270/c
LOCUS      13270 bp DNA linear PRI 30-OCT-2000
DEFINITION Homo sapiens mucolin (MCOLN1) gene, complete cds.
ACCESSION AF287270
VERSION    AF287270.1 GI:9844925
KEYWORDS
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.
REFERENCE 1 (bases 1 to 13270)
AUTHORS   Sun, M., Goldin, E., Stahl, S., Falardeau, J.L., Kennedy, J.C.,
          Acierno, J.S. Jr., Bove, C., Kaneski, C.R., Nagle, J., Bromley, M.C.,
          Colman, M., Schifmann, R. and Slaugenhaupt, S.A.
TITLE     Mucopolidosis type IV is caused by mutations in a gene encoding a
          novel transient receptor potential channel
JOURNAL   Hum. Mol. Genet. 9 (17), 2471-2478 (2000)
PUBMED    11030752
REFERENCE 2 (bases 1 to 13270)
AUTHORS   Slaugenhaupt, S.A.
TITLE     Direct Submission
JOURNAL   Submitted (13-JUL-2000) Molecular Neurogenetics, Harvard Institute
          of Human Genetics, 77 Ave. Louis Pasteur, HIM Building Room 422,
          Boston, MA 02115, USA

FEATURES             source
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     /db_xref="taxon:9606"
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6295..6403,6639..6735,6933..7032,7372..7478,7596..7745,
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LKVEFMSPCDKFAKCKPKMLQVKKLVVQLLEGSLNQLAVTRENTIAFR
HLFLGYSOGADTFRAYTREQYQALFHAVDQVLPVSLGRYAFVGGDPWING
SGALCORYYHRGHVPANDTFDIDPMVNTDCIOVDPPEPPPPDDLTLLSSSY
KNITLRFKLNVYTHIFRLTNLQSLINNEIPDCYTFSLITFDKHAHSGRIPISLE
TOAHIOECKHPSVFOHGDNSFRLLFDVVVILTCSLGFLLCARSLRGLLQNEFVGFM
WRORGVISLWSELEFVNGWYILLVTSVDLVTISGTMKIGIEAKNLASVDVCSILGT
STLLVWGVTRYLTFFHYNVILLATRLVLPVWRPCCCVAVIYLVGCGWVLGPY
HVKFRSLMSECEYDFTSLNGDDMFVTFAAQGQSSLLWLFSLYLSFISLFIYM
VLSLFTALITGTAAGTFLPGGAGAESELUQAYIAQCQDSPTSGKFRGSGSACSLICC
CGRDPSEHSLLVN"

ORIGIN
Query Match      100.0%; Score 21; DB 8; Length 13270;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 2348.47 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-4

Perfect score: 21

Sequence: 1 ctacacgtgctggaagacact 21

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST.*

1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_hc.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_est7.*
9: gb_gss1.*
10: gb_gss2.*
11: gb_gss3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	21	100.0	868	3	BI909885 603070517
C 2	18.4	87.6	225	8	CX226198 MBM01462
C 3	18.4	87.6	331	8	Z43611 HSC1GD031 n
C 4	18.4	87.6	333	8	T35364 EST83466 Hu
C 5	18.4	87.6	350	8	Z43604 HSC1GA071 n
C 6	18.4	87.6	423	8	H64102 YF57e02.r1
C 7	18.4	87.6	582	3	BP196543 BP196543
C 8	18.4	87.6	818	2	EG831705
C 9	18.4	87.6	826	2	EG913075
C 10	18.4	87.6	873	6	CD558014
C 11	18.4	87.6	878	5	BQ956452
C 12	18.4	87.6	901	5	BA433310
C 13	18.4	87.6	940	5	BQ858168
C 14	18.4	87.6	979	5	BQ918296
C 15	18.4	87.6	993	2	BG739714
C 16	18.4	87.6	1078	5	BX362786
C 17	18.4	87.6	1273	4	CR616477
C 18	18.4	87.6	1743	10	AV410283
C 19	18.4	87.6	1996	4	CR622331
C 20	18.4	87.6	3287	4	AK048278
C 21	18	85.7	901	7	CK159092
C 22	17.8	84.8	539	7	CR927955

C 23	17.8	84.8	552	3	BI775909
C 24	17.8	84.8	604	7	CN791999
C 25	17.8	84.8	660	9	BZ268910
C 26	17.8	84.8	1245	6	CA491046
C 27	17.8	84.8	1658	2	BE962361
C 28	17.4	82.9	219	3	BM293485
C 29	17.4	82.9	223	1	AI610095
C 30	17.4	82.9	271	8	N75483
C 31	17.4	82.9	276	1	AA330598
C 32	17.4	82.9	283	6	CA880163
C 33	17.4	82.9	308	1	AA879907
C 34	17.4	82.9	343	2	BB793665
C 35	17.4	82.9	364	8	H26993
C 36	17.4	82.9	375	5	BU735335
C 37	17.4	82.9	384	8	H42449
C 38	17.4	82.9	388	1	AI689890
C 39	17.4	82.9	395	2	BE690638
C 40	17.4	82.9	403	11	DE136711
C 41	17.4	82.9	409	1	AI190915
C 42	17.4	82.9	415	1	AI061413
C 43	17.4	82.9	418	1	AW172705
C 44	17.4	82.9	421	1	AA430026
C 45	17.4	82.9	421	8	N62729

ALIGNMENTS

RESULT 1
LOCUS BI909885/c
DEFINITION 603070517F1 NIH_MGC_118 Homo sapiens cDNA clone IMAGE:5219237 5',
868 bp mRNA linear EST 16-OCT-2001
mRNA sequence.
ACCESSION BI909885
VERSION BI909885.1 GI:16173182
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Homnidae; Homo.
1 (bases 1 to 868)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Life Technologies, Inc.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM1551 row: b column: 06
High quality sequence stop: 819.
Location/Qualifiers
1. .868
/organism="Homo sapiens"
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/db_xref="taxon:9606"
/clone="IMAGE:5219237"
/tissue_type="leukocyte"
/lab_host="DH10B"
/clone_lib="NIH_MGC_118"
/note="Vector: pCMV-SPORT6; Site_1: NotI; Site_2: EcoRV (destroyed); RNA source leukocytes from anonymous pool of non-activated adult donors. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.7 kb, insert size range 1.2-3.3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 027. Note:

FEATURES
source

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this is a NIH_MGC Library."

ORIGIN
Query Match      100.0%; Score 21; DB 3; Length 868;
Best Local Similarity 100.0%; Pred. No. 25;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACACT 21
    |||||
Db 122 CTCACCGTGTGGAAGACACT 102

RESULT 2
CX226198/c
LOCUS
DEFINITION
  CX226198 225 bp mRNA linear EST 29-DEC-2004
  Mus musculus hematopoietic BM-HPC5 cDNA library Mus
ACCESSION
  CX226198
VERSION
  EST.
SOURCE
  Mus musculus (house mouse)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
  Sciurognathi; Muroidea; Muridae; Murinae; Mus.
REFERENCE
  1 (bases 1 to 225)
  Williams, C., Wirta, V., Richter, K., Karlsson, C., Lundberg, J. and
  Carlsson, L.
  Expressed sequence tags of cDNA clones from a hematopoietic stem
  cell line expressing Lhx2
JOURNAL
  Unpublished (2005)
COMMENT
  Molecular Biotechnology
  Institution of Biotechnology
  Alfabova University Center, KTH-Royal Institute of Technology, 106
  91 Stockholm, Sweden
  Tel: +46855378332.
  Fax: +46855378481
  Email: cecilia.williams@biotech.kth.se
  Seq primer: M13REV.
FEATURES
  source
    1..225
    /organism="Mus musculus"
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    /strain="C57BL/6-cast"
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    /tissue_type="Bone Marrow"
    /cell_type="Hematopoietic progenitor/stem cells
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    /cell_line="hematopoietic stem cell-like cell line
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    /dev_stage="adult"
    /clone_lib="Mus Musculus hematopoietic BM-HPC5 cDNA
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    /notes="Organ: Bone Marrow; Vector: pCMVSPORT6.1;
    Preamplified custom cDNA library by Invitrogen/Reagen"

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Best Local Similarity 95.0%; Pred. No. 4.2e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACAC 20
    |||||
Db 141 CTCACCGTGTGGAAGACAC 122

RESULT 3
Z43611/c
LOCUS
DEFINITION
  Z43611 331 bp mRNA linear EST 14-NOV-1994
  HSC1GD031 normalized infant brain cDNA Homo sapiens cDNA clone
  c-1gd03, mRNA sequence.
FEATURES
  source
    1..331
    /organism="Homo sapiens"
    /mol_type="mRNA"
    /db_xref="taxon:9606"
    /clone="c-1gd03"
    /sex="Female"
    /tissue_type="total brain"
    /dev_stage="3 months old"
    /clone_lib="normalized infant brain cDNA"
    /notes="Organ: brain; Vector: lafmid BA; Site 1: HindIII;
    Site 2: NotI; sex=Female; dev stages=3 months old;
    isolate=muscular atrophy patient; tissue_type=total
    brain; total mRNA was oligo-(dT) primed and directionally
    cloned 5' -> 3' into the HindIII -> NotI sites of the
    lafmid BA vector. Clone library from B.Souares, Psychiatry
    Dept. Columbia University, USA. Normalization_method:
    Bento Soares, P.N.A.S. in press"

ORIGIN
Query Match      87.6%; Score 18.4; DB 8; Length 331;
Best Local Similarity 95.0%; Pred. No. 4.4e+02;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCACCGTGTGGAAGACACT 21
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Db 100 TCACCGTGTGGAAGACACT 81

RESULT 4
T35364/c
LOCUS
DEFINITION
  T35364 333 bp mRNA linear EST 06-SEP-1995
  EST83466 Human Brain Homo sapiens cDNA 5' end similar to None, mRNA
  sequence.
ACCESSION
  T35364
VERSION
  T35364.1 GI:617462
KEYWORDS
  EST.
SOURCE
  Homo sapiens (human)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
  Homnidae; Homo.
  1 (bases 1 to 333)
  Adams, M.D., Kerlavage, A.R., Fleischmann, R.D., Fuldner, R.A.,
  Bult, C.O., Lee, N., Kirkness, E.F., Weinstein, K.G., Gocayne, J.D.,
  White, O., Sutton, G., Blake, J.A., Brandon, R.C., Chiu, M.-W.,

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 284.928 Seconds
(without alignments)
491.207 Million cell updates/sec

Title: US-10-754-446-4

Perfect score: 21

Sequence: 1 ctcaccgtgtggaagacact 21

Scoring table:

IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: Geneseqn1980s.*

2: Geneseqn1990s.*

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4: Geneseqn2001as.*

5: Geneseqn2001bs.*

6: Geneseqn2002as.*

7: Geneseqn2002bs.*

8: Geneseqn2003as.*

9: Geneseqn2003bs.*

10: Geneseqn2003cs.*

11: Geneseqn2003ds.*

12: Geneseqn2004as.*

13: Geneseqn2004bs.*

14: Geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	21	100.0	21	14 AEB28960	Aeb28960 Human MCO
2	21	100.0	11365	4 AAK73827	Aak73827 Human imm
3	21	100.0	13270	9 ADB84283	Adb84283 Human muc
4	21	100.0	20046	4 AAK73826	Aak73826 Human imm
5	20	95.2	20	9 ADB84282	Adb84282 Human muc
6	18.4	87.6	1619	4 AAI61022	Aai61022 Human pol
7	18.4	87.6	1740	6 AAI17100	Aai17100 Human TRP
8	18.4	87.6	1740	6 ABL40755	Abi40755 Human TLC
9	18.4	87.6	1741	4 AAI59236	Aai59236 Human pol
10	18.4	87.6	1743	9 ADB84284	Adb84284 Human muc
11	18.4	87.6	1743	14 ADV66231	Adv66231 TRP-like
12	18.4	87.6	2051	13 ADP25150	Adp25150 PRO polyp
13	18.4	87.6	2052	4 AAF81753	Aaf81753 Human mem
14	18.4	87.6	2092	6 ABL90358	Abi90358 Human pol
15	18.4	87.6	2094	3 AAA39067	Aaa39067 Human sec
16	18.4	87.6	2095	6 AAI71699	Aai71699 Human TRP
17	18.4	87.6	2095	6 ABL40754	Abi40754 Human TLC
18	18.4	87.6	2095	14 ADV66229	Adv66229 TRP-like
19	18.4	87.6	2140	12 ADQ24925	Adq24925 Human sof

C	20	17.4	82.9	370	5	ABA12621	Aba12621 Human ner
C	21	17.4	82.9	421	6	ABN94509	Abn94509 Gene #100
C	22	17.4	82.9	1766	3	AAA78381	Aaa78381 Human sec
C	23	17.4	82.9	1766	8	ADA39925	Ada39925 Human sec
C	24	17.4	82.9	1766	8	ACC50509	Acc50509 Human sec
C	25	17.4	82.9	1766	8	ABZ71282	Abz71282 Secreted
C	26	17.4	82.9	1766	9	ADB91188	Adb91188 Human sec
C	27	17.4	82.9	1766	10	ADC73569	Adc73569 Human sec
C	28	17.4	82.9	1766	10	ADA56112	Ada56112 Gene enco
C	29	17.4	82.9	2530	14	AEA47619	Aea47619 Nucleotid
C	30	17.4	82.9	2750	6	ABL92130	Abi92130 Human Tum
C	31	17.4	82.9	2750	10	ABX72055	Abx72055 DNA enco
C	32	17.4	82.9	3258	4	AAH42269	Aah42269 Nucleotid
C	33	17.4	82.9	3263	10	ADA53072	Ada53072 Human cod
C	34	17.4	82.9	3272	10	ADE31397	Ade31397 Human dia
C	35	17.4	82.9	3351	11	ACN44967	Actn44967 Human mrn
C	36	17.4	82.9	125534	11	ACNA4966	Actna4966 Human gen
C	37	17	81.0	2092	5	AAS72274	Aas72274 DNA enco
C	38	16.8	80.0	102	10	ADG25870	Adg25870 INPIONCHO
C	39	16.8	80.0	297	8	ABX43933	Abx43933 Bovine ES
C	40	16.8	80.0	399	8	ABX45303	Abx45303 Bovine ES
C	41	16.8	80.0	433	8	ABX39870	Abx39870 Bovine ES
C	42	16.8	80.0	472	6	ABN73626	Abn73626 Bovine em
C	43	16.8	80.0	586	10	ADD34724	Add34724 Mouse mit
C	44	16.8	80.0	594	12	ACH77455	Ach77455 Human gen
C	45	16.8	80.0	2988	8	ACA57290	Aca57290 Human adi

ALIGNMENTS

RESULT 1

AEB28960

ID AEB28960 standard; DNA; 21 BP.

AC AEB28960;

XX 22-SEP-2005 (first entry)

DE Human MCOLN1 gene PCR primer PRI R2 SEQ ID NO:4.

KW mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; PCR;
KW primer; ss.

OS Homo sapiens.

PN US2005153300-A1.

PD 14-JUL-2005.

PF 09-JAN-2004; 2004US-00754446.

PR 09-JAN-2004; 2004US-00754446.

XX (QUES-) QUEST DIAGNOSTICS INC.

XX Sun W, Hantash F;

XX WPI; 2005-521160/53.

XX Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by
amplifying the nucleic acid, detecting amplified product with labeled
oligonucleotide probes via a change in fluorescence which indicates the
presence of an ML IV mutant.

PS Claim 1; SEQ ID NO 4; 15pp; English.

XX The invention relates to a method (M1) for detecting the presence of
mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1)
involves contacting the nucleic acid with oligonucleotide primers and
probes, conducting amplification by temperature cycling and monitoring
the accumulation of amplified nucleic acid by detecting an increase in
donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
CC described: (1) an oligonucleotide (ON1) comprising a sequence
CC complementary to the coding or non-coding strand of the mucolin-1
CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
CC sequences, comprising ON1. (M1) is useful for determining the presence of
CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
CC is useful for genetic counseling of individuals at risk for ML IV or at
CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
CC on chromosome 19p13. The present sequence represents a PCR primer for the
CC MCOLN1 gene, which is used in the exemplification of the present
CC invention.

XX
SQ Sequence 21 BP; 5 A; 7 C; 5 G; 4 T; 0 U; 0 Other;

Query Match 100.0%; Score 21; DB 14; Length 21;

Best Local Similarity 100.0%; Pred. No. 2;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCACCGTGTGGAGACACT 21

Db 1 CTCACCGTGTGGAGACACT 21

RESULT 2

AAK73827/c

ID AAK73827 standard; DNA; 11365 BP.

XX AC AAK73827;

XX DT 07-NOV-2001 (first entry)

XX DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:28639.

XX KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;

XX KW Cytostatic; gene therapy; vaccine; metastasis; ds.

XX OS Homo sapiens.

XX PN WO200157182-A2.

XX PD 09-AUG-2001.

XX PF 17-JAN-2001; 2001WO-US001354.

XX PR 31-JAN-2000; 2000US-0179065P.

XX PR 04-FEB-2000; 2000US-0180628P.

XX PR 24-FEB-2000; 2000US-0184664P.

XX PR 02-MAR-2000; 2000US-0186350P.

XX PR 16-MAR-2000; 2000US-0189874P.

XX PR 17-MAR-2000; 2000US-0190076P.

XX PR 18-APR-2000; 2000US-0198123P.

XX PR 19-MAY-2000; 2000US-0205515P.

XX PR 07-JUN-2000; 2000US-0209467P.

XX PR 28-JUN-2000; 2000US-0214886P.

XX PR 30-JUN-2000; 2000US-0215135P.

XX PR 07-JUL-2000; 2000US-0216647P.

XX PR 07-JUL-2000; 2000US-0216880P.

XX PR 11-JUL-2000; 2000US-0217487P.

XX PR 11-JUL-2000; 2000US-0217496P.

XX PR 14-JUL-2000; 2000US-0218290P.

XX PR 26-JUL-2000; 2000US-0220963P.

XX PR 26-JUL-2000; 2000US-0220964P.

XX PR 14-AUG-2000; 2000US-0224518P.

XX PR 14-AUG-2000; 2000US-0224519P.

XX PR 14-AUG-2000; 2000US-0225213P.

XX PR 14-AUG-2000; 2000US-0225214P.

XX PR 14-AUG-2000; 2000US-0225266P.

XX PR 14-AUG-2000; 2000US-0225267P.

XX PR 14-AUG-2000; 2000US-0225268P.

XX PR 14-AUG-2000; 2000US-0225270P.

XX PR 14-AUG-2000; 2000US-0225447P.

XX PR 14-AUG-2000; 2000US-0225757P.

XX PR 14-AUG-2000; 2000US-0225758P.

PR 14-AUG-2000; 2000US-0225759P.
PR 18-AUG-2000; 2000US-0226279P.
PR 22-AUG-2000; 2000US-0226681P.
PR 22-AUG-2000; 2000US-0226688P.
PR 22-AUG-2000; 2000US-0227182P.
PR 23-AUG-2000; 2000US-0227009P.
PR 30-AUG-2000; 2000US-0228924P.
PR 01-SEP-2000; 2000US-0229287P.
PR 01-SEP-2000; 2000US-0229343P.
PR 01-SEP-2000; 2000US-0229344P.
PR 01-SEP-2000; 2000US-0229345P.
PR 05-SEP-2000; 2000US-0229509P.
PR 05-SEP-2000; 2000US-0229513P.
PR 06-SEP-2000; 2000US-0230437P.
PR 06-SEP-2000; 2000US-0230438P.
PR 08-SEP-2000; 2000US-0231422P.
PR 08-SEP-2000; 2000US-0231423P.
PR 08-SEP-2000; 2000US-0231433P.
PR 08-SEP-2000; 2000US-0231413P.
PR 08-SEP-2000; 2000US-0231414P.
PR 08-SEP-2000; 2000US-0232080P.
PR 12-SEP-2000; 2000US-0232081P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
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C 3	20	95.2	20	3	US-09-851-494B-11	Sequence 11, Appli	
C 4	18.4	87.6	25	10	US-11-060-756-194946	Sequence 194946,	
C 5	18.4	87.6	1400	10	US-11-060-756-2850	Sequence 2850, A	
C 6	18.4	87.6	1400	10	US-11-060-756-7122	Sequence 7122, A	
C 7	18.4	87.6	1740	3	US-09-828-466-3	Sequence 3, Appli	
C 8	18.4	87.6	1740	5	US-10-103-458-3	Sequence 3, Appli	
C 9	18.4	87.6	1743	8	US-10-782-695-6	Sequence 6, Appli	
C 10	18.4	87.6	2051	3	US-09-851-494B-2	Sequence 2, Appli	
C 11	18.4	87.6	2052	3	US-09-855-529-50	Sequence 50, Appl	
C 12	18.4	87.6	2052	3	US-09-969-680A-50	Sequence 50, Appl	
C 13	18.4	87.6	2052	10	US-11-048-692-50	Sequence 50, Appl	
C 14	18.4	87.6	2032	6	US-10-264-237-920	Sequence 50, Appl	
C 15	18.4	87.6	2094	3	US-09-820-893-26	Sequence 26, Appl	
C 16	18.4	87.6	2094	7	US-10-607-565-26	Sequence 26, Appl	
C 17	18.4	87.6	2095	3	US-09-828-466-1	Sequence 1, Appli	
C 18	18.4	87.6	2095	5	US-10-103-458-1	Sequence 1, Appli	
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C 20	18.4	87.6	2120	8	US-10-723-860-7745	Sequence 7745, Ap	
C 21	17.4	82.9	421	3	US-09-880-107-1007	Sequence 1007, Ap	
C 22	17.4	82.9	644	5	US-10-027-632-193464	Sequence 193464,	
C 23	17.4	82.9	644	6	US-10-027-632-193464	Sequence 193464,	

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; SEQ ID NO 1
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; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-851-494B-1

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DB 7037 CTCACCGTGTGGAAGACACT 7017

RESULT 3

US-09-851-494B-11
; Sequence 11, Application US/09851494B
; Publication No. US20030064363A1
; GENERAL INFORMATION:
; APPLICANT: ML4 Foundation
; APPLICANT: Goldin, Ehud
; APPLICANT: Slangenhuysen, Susan A.
; APPLICANT: Sun, Mei
; APPLICANT: Acierno, James S.
; TITLE OF INVENTION: A Gene Encoding A New TRP Channel is Mutated in Mucopolidosis IV
; FILE REFERENCE: 3394/1H557US1
; CURRENT APPLICATION NUMBER: US/09/851,494B
; CURRENT FILING DATE: 2002-07-12
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 11
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: PCR primer
US-09-851-494B-11

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Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DB 1 CTCACCGTGTGGAAGACAC 20

RESULT 4

US-11-060-756-194946/c
; Sequence 194946, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 194946
; LENGTH: 25
; TYPE: DNA
; ORGANISM: probe
US-11-060-756-194946

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Best Local Similarity 95.0%; Pred. No. 24;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 22 TCTCCGTGTGGAAGACACT 3

RESULT 5

US-11-060-756-2850/c
; Sequence 2850, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2850
; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-2850

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Best Local Similarity 95.0%; Pred. No. 23;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCACCGTGTGGAAGACACT 21
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DB 355 TCTCCGTGTGGAAGACACT 336

RESULT 6

US-11-060-756-7122/c
; Sequence 7122, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7122
; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-7122

Query Match 87.6%; Score 18.4; DB 10; Length 1400;
Best Local Similarity 95.0%; Pred. No. 23;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCACCGTGTGGAAGACACT 21
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DB 355 TCTCCGTGTGGAAGACACT 336

RESULT 7

US-09-828-466-3/c
; Sequence 3, Application US/09828466
; Patent No. US20020035056A1
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Silos-Santiago, Immaculada
; TITLE OF INVENTION: 54420, A NOVEL HUMAN CALCIUM CHANNEL
; FILE REFERENCE: WNI-125CP
; CURRENT APPLICATION NUMBER: US/09/828,466

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	15.4	73.3	1238	6	US-11-083-784-151656
5	15.4	73.3	1238	6	US-10-821-234-1
6	15.4	73.3	1244	7	US-11-000-688-733
7	15.4	73.3	6990	7	US-11-000-688-609
8	15.4	73.3	175673	7	US-11-121-086-55
9	15.2	72.4	21	6	US-10-310-914A-1157477
10	15.2	72.4	579	7	US-11-128-061-342
11	15.2	72.4	579	7	US-11-128-061-3984
12	15.2	72.4	866	6	US-10-750-185-61786
13	15.2	72.4	866	6	US-10-750-623-61786
14	15.2	72.4	1008	6	US-10-750-185-54439
15	15.2	72.4	1007	6	US-10-750-623-54439
16	15.2	72.4	1077	6	US-10-517-939-369
17	15.2	72.4	1307	6	US-10-750-185-27270
18	15.2	72.4	1307	6	US-10-750-623-27270
19	15.2	72.4	1350	7	US-11-055-822-289
20	15.2	72.4	3272	6	US-10-750-185-24646
21	15.2	72.4	3272	6	US-10-750-623-24646
22	15.2	72.4	4730	7	US-11-128-061-520
23	15.2	72.4	5595	6	US-10-955-054A-11

c 24	15.2	72.4	154452	7	US-11-121-086-74	Sequence 74, Appl
c 25	15	71.4	201	6	US-10-995-561-26690	Sequence 26690, A
c 26	15	71.4	201	6	US-10-995-561-73444	Sequence 73444, A
c 27	15	71.4	14082	6	US-10-995-561-13445	Sequence 13445, A
c 28	15	71.4	305312	6	US-10-995-561-13236	Sequence 13236, A
c 29	14.8	70.5	19	8	US-11-101-244-285326	Sequence 285326, A
c 30	14.8	70.5	19	9	US-11-083-784-285326	Sequence 285326, A
c 31	14.8	70.5	25	7	US-11-121-849-182614	Sequence 182614, A
c 32	14.8	70.5	201	6	US-10-995-561-23969	Sequence 23969, A
c 33	14.8	70.5	327	7	US-11-000-688-786	Sequence 786, App
c 34	14.8	70.5	1105	6	US-11-000-463-581	Sequence 581, App
c 35	14.8	70.5	1296	6	US-10-525-710-23	Sequence 23, Appl
c 36	14.8	70.5	1388	7	US-11-128-061-969	Sequence 969, App
c 37	14.8	70.5	1481	7	US-11-000-688-788	Sequence 788, App
c 38	14.8	70.5	1646	6	US-10-750-185-55975	Sequence 55975, A
c 39	14.8	70.5	1646	6	US-10-750-623-55975	Sequence 55975, A
c 40	14.8	70.5	2244	7	US-11-128-061-528	Sequence 528, App
c 41	14.8	70.5	98345	7	US-11-112-908-36	Sequence 36, Appl
c 42	14.8	70.5	127340	7	US-11-112-908-35	Sequence 35, Appl
c 43	14.8	70.5	207835	7	US-11-121-086-39	Sequence 39, Appl
c 44	14.8	70.5	207835	7	US-11-121-086-40	Sequence 40, Appl
c 45	14.6	69.5	25	7	US-11-136-527-319963	Sequence 319963, A

ALIGNMENTS

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; Sequence 2907, Application US/11124368A
; Publication No. US20050287559A1
; GENERAL INFORMATION:
; APPLICANT: Michele Cargill
; APPLICANT: James J. Devlin
; APPLICANT: May Luke
; TITLE OF INVENTION: Genetic Polymorphisms Associated with
; TITLE OF INVENTION: Vascular Diseases, Methods of Detection and Uses Thereof
; FILE REFERENCE: CL001524
; CURRENT APPLICATION NUMBER: US/11/124,368A
; PRIOR FILING DATE: 2005-05-09
; PRIOR FILING DATE: 2004-05-07
; PRIOR FILING DATE: 2004-05-07
; PRIOR FILING DATE: 2004-11-09
; NUMBER OF SEQ ID NOS: 21112
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2907
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; ORGANISM: Homo sapiens
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; OTHER INFORMATION: n = A,T,C or G
US-11-124-368A-2907

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; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; TITLE OF INVENTION: CARDIOVASCULAR DISORDERS AND DRUG RESPONSE, METHODS OF
; TITLE OF INVENTION: DETECTION AND USES THEREOF
; FILE REFERENCE: CL001559
; CURRENT APPLICATION NUMBER: US/10/995,561
; CURRENT FILING DATE: 2004-11-24
; NUMBER OF SEQ ID NOS: 85702
; SOFTWARE: FastSeq for Windows Version 4.0
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; TYPE: DNA
; ORGANISM: Homo sapiens
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Db 17401 TCACCGTGTGTGGGAAGAAC 17383

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; Publication No. US20050246794A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 134990S
; CURRENT APPLICATION NUMBER: US/11/101,244

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; CURRENT FILING DATE: 2005-04-07
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
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; SOFTWARE: Proprietary
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; TYPE: RNA
; ORGANISM: Homo sapiens
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; Publication No. US20050245475A1
; GENERAL INFORMATION:
; APPLICANT: Dharmacon, Inc.
; APPLICANT: Khvorova, Anastasia
; APPLICANT: Reynolds, Angela
; APPLICANT: Leake, Devin
; APPLICANT: Marshall, William
; APPLICANT: Scaringe, Stephen
; TITLE OF INVENTION: Functional and Hyperfunctional siRNA
; FILE REFERENCE: 134990S
; CURRENT APPLICATION NUMBER: US/11/083,784
; CURRENT FILING DATE: 2005-03-18
; PRIOR APPLICATION NUMBER: US/10/714,333
; PRIOR FILING DATE: 2003-11-14
; PRIOR APPLICATION NUMBER: 60/502,050
; PRIOR FILING DATE: 2003-09-10
; PRIOR APPLICATION NUMBER: 60/426,137
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; ORGANISM: Homo sapiens
US-11-083-784-151656

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Best Local Similarity 76.5%; Pred. No. 1.3e+02;
Matches 13; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

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Db 1 UCAACGUGUGGAAGAC 17

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; Publication No. US20050255114A1
; GENERAL INFORMATION:
; APPLICANT: Labat, Ivan
; APPLICANT: Stache-Grain, Birgit
; APPLICANT: Andarmani, Susan
; APPLICANT: Tang, Y. Tom
; TITLE OF INVENTION: Methods for Diagnosis and Treatment of Preeclampsia
; FILE REFERENCE: 821A
; CURRENT APPLICATION NUMBER: US/10/821,234
; CURRENT FILING DATE: 2004-04-07

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5: /cgn2_6/ptodata/1/ina/H/COMB.seq.*
6: /cgn2_6/ptodata/1/ina/PC/COMB.seq.*
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8: /cgn2_6/ptodata/1/ina/RE/COMB.seq.*
9: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	21	100.0	15353	3	US-09-949-016-13454
C 3	18.4	87.6	2051	3	US-09-949-016-1712
C 4	16.2	77.1	25	3	US-09-396-196G-50417
C 5	16.2	77.1	471	3	US-09-270-767-1970
C 6	16.2	77.1	471	3	US-09-270-767-17252
C 7	16.2	77.1	601	3	US-09-949-016-178570
C 8	16.2	77.1	601	3	US-09-949-016-178571
C 9	16.2	77.1	797	3	US-09-270-767-7266
C 10	16.2	77.1	797	3	US-09-270-767-22548
C 11	16.2	77.1	3385	3	US-09-509-800-1
C 12	16.2	77.1	12470	3	US-09-949-002-847
C 13	16.2	77.1	17896	3	US-09-949-016-16867
C 14	16.2	77.1	32665	3	US-09-949-016-14098
C 15	16.2	77.1	52667	3	US-09-949-016-12019
C 16	16.2	77.1	141560	3	US-09-949-016-16476
C 17	15.8	75.2	25	3	US-09-396-196G-50406
C 18	15.8	75.2	659	3	US-09-533-559-6889
C 19	15.8	75.2	1293	3	US-09-902-540-5426
C 20	15.8	75.2	24154	3	US-09-949-016-16374
C 21	15.8	75.2	27465	3	US-09-949-016-16561
C 22	15.8	75.2	34953	3	US-09-902-540-1263
C 23	15.4	73.3	358	3	US-09-513-999C-3422
C 24	15.4	73.3	601	3	US-09-949-016-51396

25	15.4	73.3	624	3	US-09-513-999C-2084	Sequence 2084, Ap
26	15.4	73.3	717	3	US-09-949-002-92	Sequence 92, Appl
27	15.4	73.3	717	3	US-09-949-002-127	Sequence 127, App
28	15.4	73.3	1037	2	US-08-240-124-1	Sequence 1, Appli
29	15.4	73.3	1037	2	US-08-453-943-1	Sequence 1, Appli
30	15.4	73.3	1037	2	US-09-057-121-1	Sequence 1, Appli
31	15.4	73.3	1037	2	US-09-358-734-1	Sequence 1, Appli
32	15.4	73.3	1037	3	US-09-904-954-1	Sequence 1, Appli
33	15.4	73.3	1070	2	US-08-299-567-8	Sequence 8, Appli
34	15.4	73.3	1092	2	US-08-646-590B-35	Sequence 35, Appl
35	15.4	73.3	1092	3	US-09-412-184-35	Sequence 35, Appl
36	15.4	73.3	1212	3	US-09-949-016-1491	Sequence 1491, Ap
37	15.4	73.3	1374	3	US-09-902-540-4675	Sequence 4675, Ap
38	15.4	73.3	1533	3	US-09-711-164-194	Sequence 194, App
39	15.4	73.3	2646	3	US-09-489-039A-4832	Sequence 4832, Ap
C 40	15.4	73.3	6503	3	US-09-404-650-12	Sequence 12, Appl
C 41	15.4	73.3	6503	3	US-09-935-541-12	Sequence 12, Appl
C 42	15.4	73.3	6503	3	US-10-425-800-12	Sequence 12, Appl
C 43	15.4	73.3	6816	3	US-09-404-650-1	Sequence 1, Appli
C 44	15.4	73.3	6816	3	US-09-935-541-1	Sequence 1, Appli
C 45	15.4	73.3	6816	3	US-10-425-800-1	Sequence 1, Appli

ALIGNMENTS

RESULT 1

US-09-949-016-59085/c
; Sequence 59085, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 59085
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-59085

Query Match 100.0%; Score 21; DB 3; Length 601;

Best Local Similarity 100.0%; Pred. No. 0.99; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCACCGTGTGGAAGACACT 21
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Db 504 CTCACCGTGTGGAAGACACT 484

RESULT 2

US-09-949-016-13454/c
; Sequence 13454, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20

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; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13454
; LENGTH: 15353
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13454

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Best Local Similarity 100.0%; Pred. No. 1.3;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACACT 21
Db 7637 CTCACCGTGTGGAAGACACT 7617

RESULT 3
US-09-949-016-1712/c
; Sequence 1712, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; FILE REFERENCE: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1712
; LENGTH: 2051
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-1712

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Best Local Similarity 95.0%; Pred. No. 20;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCACCGTGTGGAAGACACT 21
Db 1006 TCACCGTGTGGAAGACACT 987

RESULT 4
US-09-396-196G-50417
; Sequence 50417, Application US/09396196G
; Patent No. 6821724
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; APPLICANT: David Mack
; APPLICANT: David Lockhart
; APPLICANT: Affymetrix, Inc.
; TITLE OF INVENTION: Methods of Genetic Analysis
; FILE REFERENCE: 3101.1
; CURRENT FILING DATE: 1999-09-15
; PRIOR APPLICATION NUMBER: US/09/396,196G
; CURRENT FILING DATE: 1999-09-15
; PRIOR APPLICATION NUMBER: 60/100,678
; PRIOR FILING DATE: 1998-09-17
; NUMBER OF SEQ ID NOS: 127806
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 50417
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; LENGTH: 25
; TYPE: DNA
; ORGANISM: mus musculus
US-09-396-196G-50417

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Best Local Similarity 85.7%; Pred. No. 1.7e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACACT 21
Db 4 CACACCGTGTGGAAGACACT 24

RESULT 5
US-09-270-767-1970/c
; Sequence 1970, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1970
; LENGTH: 471
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-1970

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Best Local Similarity 85.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACACT 21
Db 451 CTCACCGTGTGGAAGACACT 431

RESULT 6
US-09-270-767-17252/c
; Sequence 17252, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 17252
; LENGTH: 471
; TYPE: DNA
; ORGANISM: Drosophila melanogaster
US-09-270-767-17252

Query Match      77.1%; Score 16.2; DB 3; Length 471;
Best Local Similarity 85.7%; Pred. No. 2.1e+02;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CTCACCGTGTGGAAGACACT 21
Db 451 CTCACCGTGTGGAAGACACT 431

RESULT 7
US-09-949-016-178570
; Sequence 178570, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
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GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 982.24 Seconds
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1157.425 Million cell updates/sec

Title: US-10-754-446-3

Perfect score: 20

Sequence: 1 cttgctctgtgccaggct 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 5883141 seqs, 28421725653 residues

Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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3: gb_env.*
4: gb_on.*
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14: gb_htg.*
15: gb_pl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	20	100.0	51	6 AX163311	Sequence
C 2	20	100.0	51	6 AX199337	Sequence
C 3	20	100.0	51	6 AX199338	Sequence
C 4	20	100.0	71	6 AX522607	Sequence
C 5	20	100.0	77	6 AX523183	Sequence
C 6	20	100.0	98	6 BD039678	Sequence
C 7	20	100.0	98	6 AX904145	Sequence
C 8	20	100.0	102	6 BD037012	Sequence
C 9	20	100.0	102	6 AX901479	Sequence
C 10	20	100.0	107	6 CQ685626	Sequence
C 11	20	100.0	108	6 CQ694139	Sequence
C 12	20	100.0	114	6 BD044993	Sequence
C 13	20	100.0	114	6 AX909460	Sequence
C 14	20	100.0	117	6 BD037394	Sequence
C 15	20	100.0	117	6 BD044139	Sequence
C 16	20	100.0	117	6 AX901861	Sequence
C 17	20	100.0	117	6 AX908606	Sequence
C 18	20	100.0	128	6 BD034714	Sequence

19	20	100.0	128	6 AX899181	Sequence
C 20	20	100.0	129	6 CQ708055	Sequence
21	20	100.0	145	6 BD035848	Sequence
22	20	100.0	145	6 BD045580	Sequence
C 23	20	100.0	145	6 BD118446	Sequence
C 24	20	100.0	145	6 AR422893	Sequence
25	20	100.0	145	6 AX900315	Sequence
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C 27	20	100.0	145	6 AX983587	Sequence
C 28	20	100.0	147	6 CQ686625	Sequence
29	20	100.0	151	6 BD043797	Sequence
C 30	20	100.0	151	6 AX908264	Sequence
C 31	20	100.0	152	6 BD115706	Sequence
C 32	20	100.0	152	6 BD117639	Sequence
C 33	20	100.0	152	6 AR420153	Sequence
C 34	20	100.0	152	6 AR422086	Sequence
C 35	20	100.0	152	6 AX980847	Sequence
C 36	20	100.0	152	6 AX982780	Sequence
37	20	100.0	154	10 AB134303	Sequence
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C 41	20	100.0	158	6 BD116463	Sequence
C 42	20	100.0	158	6 AR420191	Sequence
C 43	20	100.0	158	6 AR420910	Sequence
C 44	20	100.0	158	6 AX980885	Sequence
C 45	20	100.0	158	6 AX981604	Sequence

ALIGNMENTS

RESULT 1
AX163311/c
LOCUS AX163311 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 6639 from Patent WO0140521.
ACCESSION AX163311
VERSION AX163311.1 GI:14544642
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0140521-A 6639 07-JUN-2001;
FEATURES Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
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misc_feature 26
/note="1 of 2 allelic variants (6640 is other entry)
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Best Local Similarity 100.0%; Pred.No.11;
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Db 37 CTTGCTCTGTGCCAGGCT 18

RESULT 2
AX199337/c
LOCUS AX199337 51 bp DNA linear PAT 29-AUG-2001
DEFINITION Sequence 267 from Patent WO0151670.

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ACCESSION AX199337
VERSION AX199337.1 GI:15389718
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
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          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
          Hominiidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.D.
TITLE Nucleic acids containing single nucleotide polymorphisms and
        methods of use thereof
JOURNAL Patent: WO 0151670-A 267 19-JUL-2001;
        Curagen Corporation (US)
FEATURES
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              /mol_type="unassigned DNA"
              /db_xref="taxon:9606"
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Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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Db 49 CTTGCTCTGTGCCAGGCT 30

RESULT 3
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LOCUS AX199338
DEFINITION Sequence 268 from Patent WO0151670.
ACCESSION AX199338
VERSION AX199338.1 GI:15389719
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
          Hominiidae; Homo.
REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.D.
TITLE Nucleic acids containing single nucleotide polymorphisms and
        methods of use thereof
JOURNAL Patent: WO 0151670-A 268 19-JUL-2001;
        Curagen Corporation (US)
FEATURES
  source      1..51
              /organism="Homo sapiens"
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Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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RESULT 4
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LOCUS AX522607
DEFINITION Sequence 277 from Patent WO02064731.
ACCESSION AX522607.1 GI:24411561
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
          Hominiidae; Homo.
REFERENCE 1
AUTHORS Telerman,A., Anson,R., Tuijnder,M. and Susini,L.
TITLE Sequences involved in phenomena of tumour suppression, tumour
        reversion, apoptosis and/or virus resistance and their use as
        medicines
JOURNAL Patent: WO 02064731-A 277 22-AUG-2002;
        Molecular Engines Laboratories (FR)
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Db 11 CTTGCTCTGTGCCAGGCT 30

RESULT 5
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LOCUS AX523183
DEFINITION Sequence 853 from Patent WO02064731.
ACCESSION AX523183
VERSION AX523183.1 GI:24412137
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
          Hominiidae; Homo.
REFERENCE 1
AUTHORS Telerman,A., Anson,R., Tuijnder,M. and Susini,L.
TITLE Sequences involved in phenomena of tumour suppression, tumour
        reversion, apoptosis and/or virus resistance and their use as
        medicines
JOURNAL Patent: WO 02064731-A 853 22-AUG-2002;
        Molecular Engines Laboratories (FR)
FEATURES
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              /mol_type="unassigned DNA"
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Best Local Similarity 100.0%; Pred. No. 11;
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    |||||
Db 39 CTTGCTCTGTGCCAGGCT 20

RESULT 6
BD039678/c
LOCUS BD039678
DEFINITION Sequence tag and encoded human protein.
ACCESSION BD039678
VERSION BD039678.1 GI:22581420

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GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 2236.64 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-3

Perfect score: 20

Sequence: 1 cttgctgtgtgccaggct 20

Scoring table: IDENTITY_NUC

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Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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7: gb_est6.*

8: gb_est7.*

9: gb_gss1.*

10: gb_gss2.*

11: gb_gss3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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c 3	20	100.0	46	9	AZ576121 AST-T32E0
c 4	20	100.0	49	5	C20875 HUMS000494
c 5	20	100.0	50	1	AI708455 as97e06.x
c 6	20	100.0	50	1	AU102403 AU102403
c 7	20	100.0	52	1	AA903439 ok48h04.s
c 8	20	100.0	61	1	AI735552 at17e12.x
c 9	20	100.0	64	1	AW304187 xv61e11.x
c 10	20	100.0	68	9	AZ537585 AST-2P881
c 11	20	100.0	69	1	AA215549 zr97e12.r
c 12	20	100.0	70	1	AA604042 no44h11.s
c 13	20	100.0	82	2	BG236597 nai45g03.
c 14	20	100.0	84	9	AZ757146 ew06c09.r
c 15	20	100.0	101	1	AA262246 z826e09.s
c 16	20	100.0	102	6	CD640446
c 17	20	100.0	106	2	BF875905 RC3-ET013
c 18	20	100.0	108	2	BE141837 IL5-HT011
c 19	20	100.0	108	8	T58114 yb26d03.r1
c 20	20	100.0	111	2	BE246735 TCBAPI044
c 21	20	100.0	111	9	B66475 CIT-HSP-201
c 22	20	100.0	113	1	AW780390 ho05a12.x

c 23	20	100.0	116	3	BQ372956
c 24	20	100.0	116	6	CD521979
c 25	20	100.0	116	8	N83678
c 26	20	100.0	118	1	AA004455
c 27	20	100.0	118	8	H57636
c 28	20	100.0	120	1	AA007673
c 29	20	100.0	120	1	AA013351
c 30	20	100.0	120	1	AA367235
c 31	20	100.0	122	2	BG956764
c 32	20	100.0	122	2	BG957876
c 33	20	100.0	122	2	BG957892
c 34	20	100.0	123	7	CK854800
c 35	20	100.0	123	8	R92902
c 36	20	100.0	125	2	BF820510
c 37	20	100.0	125	8	H71447
c 38	20	100.0	125	8	H73044
c 39	20	100.0	125	8	H73847
c 40	20	100.0	126	1	AA579709
c 41	20	100.0	126	9	AQ201947
c 42	20	100.0	128	1	AA328830
c 43	20	100.0	128	8	H67228
c 44	20	100.0	128	8	T83695
c 45	20	100.0	130	1	AA659815

ALIGNMENTS

RESULT 1	AA054107/c	38 bp	mrna	linear	EST 13-SEP-1996
LOCUS	zf50f06.r1	Soares retina N2b4HR	Homo sapiens	cDNA clone	
DEFINITION	IMAGE:380387 5', similar to gb:X77738_rnal BAND 3 ANION TRANSPORT PROTEIN (HUMAN) ; , mRNA sequence.				
ACCESSION	AA054107				
VERSION	AA054107.1	GI:1545030			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.				
AUTHORS	Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Rifkin,L., Rohlfs,T., Soares,M., Tan,F., Trevasaki,E., Waterston,R., Williamson,A., Wohlmann,P. and Wilson,R.				
TITLE	The WashU-Merck EST Project				
JOURNAL	Unpublished (1995)				
COMMENT	Contact: Wilson RK Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu This clone is available royalty-free through LLNL ; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Trace considered overall poor quality Seq primer: -28M13 rev2 from Amersham High quality sequence stop: 1. Location/Qualifiers 1. .38 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="GDB:1288644" /db_xref="taxon:9606" /clone="IMAGE:380387" /sex="male" /tissue type="retina" /dev stage="55 year old" /lab_host="DH10B (ampicillin resistant)" /clone_lib="Soares retina N2b4HR"				

/note="Organ: eye; Vector: pT73D (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' TGTTACCAATCTGAAGTGGAGCGCCGCTTTTCTTTTCTTTT 3'], double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). The retinas were obtained from a 55 year old Caucasian and total cellular poly(A)+ RNA was extracted 6 hrs after their removal. The retina RNA was kindly provided by Roderick R. McInnes M.D. Ph.D. from the University of Toronto. Library constructed by Bento Soares and M.Fatima Bonaldo. "

ORIGIN

Query Match 100.0%; Score 20; DB 1; Length 38;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
 |||||
 Db 28 CTTGCTCTGTGGCCAGGCT 9

RESULT 2

AA911358
 LOCUS
 DEFINITION
 o676d10.61 NCI CGAP Lu5 Homo sapiens cDNA clone IMAGE:1417555 3' similar to gb:X01057_rnal INTERLEUKIN-2 RECEPTOR ALPHA CHAIN PRECURSOR (HUMAN); contains element MER22 repetitive element ;, mRNA sequence.

ACCESSION

AA911358.1 GI:3050722

VERSION

EST.

KEYWORDS

Source

ORGANISM

Homo sapiens (human)
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.
 1 (bases 1 to 39)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 Unpublished (1997)
 Contact: Robert Strausberg, Ph.D.
 Email: cgaps-rc@mail.nih.gov
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 DNA Library Arrayed by: Greg Lennon, Ph.D.
 Cloning by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html

FEATURES

source

1. .39
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue="IMAG:1417555"
 /tissue_type="carcinoid"
 /lab_host="DH10B"
 /clone_lib="NCI CGAP Lu5"
 /note="Organ: lung; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from neuroendocrine lung carcinoma, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and

ORIGIN

Query Match 100.0%; Score 20; DB 1; Length 39;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
 |||||
 Db 8 CTTGCTCTGTGGCCAGGCT 27

RESULT 3

AAZ576121

LOCUS

DEFINITION

sapiens genomic 5', genomic survey sequence.

AAZ576121

AAZ576121.1 GI:11562432

GSS.

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Homiidae; Homo.

1 (bases 1 to 46)

Henkel, G., Liyanage, M., Pratt, E., Huang, D., Riley, M., Bernardino, A., Durick, K. and Pollok, B.

Exon-trap tags from a T47D GenomeScreen(TM) Library

Unpublished (2000)

Contact: Greg Henkel

Gene Expression

Aurora Biosciences Corp.

11010 Torreyana Road, San Diego, CA 92121, USA

Tel: 8584048436

Fax: 8584046719

Email: henkel@aurorabio.com

Pools of cells were isolated from a GenomeScreen(TM) library. The library of cells was generated by retroviral integration of a gene tagging element consisting of: 1) A promoterless beta-lactamase

proceeded by a splice acceptor as a reporter for gene expression;

2) A promoter driving neomycin resistance followed by a splice

donor to trap downstream exons. 3' RACE from neomycin gene was

performed using total RNA from isolated pools. Output was shotgun

cloned in pAMP-1 and used to transform DH5-alpha competent

bacteria. 5' ends of reported sequences were immediately preceded

by splice donor from the trapping construct.

Class: exon-trapped.

Location/Qualifiers

1. .46

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/tissue_type="Carcinoma"

/cell_type="Epithelial"

/cell_line="T47D"

/clone_lib="Genetrap T47D Human Breast Carcinoma Library"

/notes="Organ: Breast; Vector: pAMP-1; 3' RACE of total RNA

from genetrap pools; shotgun clone in pAMP-1 and used to

transform DH5-alpha competent bacteria."

ORIGIN

Query Match 100.0%; Score 20; DB 9; Length 46;
 Best Local Similarity 100.0%; Pred. No. 2.7e+02;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
 |||||
 Db 6 CTTGCTCTGTGGCCAGGCT 25

cloned into the Not I and Eco RI sites of the modified pT73 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo. "

GenCore version 5.1.6
Copyright (c) 1993 - 2006 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 271.36 Seconds
(without alignments)
491.207 Million cell updates/sec

Title: US-10-754-446-3

Perfect score: 20

Sequence: 1 cttgctctgtgtccaggct 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N Geneseq 21.*

- 1: Geneseqn1980s.*
- 2: Geneseqn1990s.*
- 3: Geneseqn2000s.*
- 4: Geneseqn2001as.*
- 5: Geneseqn2001bs.*
- 6: Geneseqn2002as.*
- 7: Geneseqn2002bs.*
- 8: Geneseqn2003as.*
- 9: Geneseqn2003bs.*
- 10: Geneseqn2003cs.*
- 11: Geneseqn2003ds.*
- 12: Geneseqn2004as.*
- 13: Geneseqn2004bs.*
- 14: Geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	14	AEB28959 Human MCO
c	2	20	100.0	24	ABV75668 Human PD1
3	20	100.0	41	AAH48082	Aah48082 Ras GTPas
c	4	20	100.0	41	AAH89487 Human pro
c	5	20	100.0	41	AAH89487 Human pro
c	6	20	100.0	41	AAH89487 Human pro
c	7	20	100.0	51	AAH89487 Human con
c	8	20	100.0	51	AAH89487 Human cod
c	9	20	100.0	51	AAH89487 Human cod
10	20	100.0	60	10	ADI20565 Oligonucle
11	20	100.0	71	8	ABZ09117 Human oli
12	20	100.0	71	10	ABZ78570 Tumour su
c	13	20	100.0	77	ABZ09693 Human oli
c	14	20	100.0	77	ABZ79146 Tumour su
15	20	100.0	82	13	ADR16680 MicroRNA
16	20	100.0	84	4	AAH89487 Human pro
17	20	100.0	84	10	ADG41734 Human res
18	20	100.0	84	11	ADI97508 Human res
c	19	20	100.0	91	AAH26410 Human gen

c	20	20	100.0	93	4	AAK65580	Aak65580 Human imm
c	21	20	100.0	94	4	AAK73464	Aak73464 Human imm
c	22	20	100.0	94	4	AAK72995	Aak72995 Human imm
c	23	20	100.0	95	4	AAK77204	Aak77204 Human imm
c	24	20	100.0	95	4	AAK77203	Aak77203 Human imm
c	25	20	100.0	95	4	AAK70695	Aak70695 Human imm
c	26	20	100.0	98	3	AAK15933	Aak15933 Human sec
c	27	20	100.0	98	4	AAK36698	Aak36698 Human car
c	28	20	100.0	98	4	AAK71993	Aak71993 Human imm
c	29	20	100.0	98	10	AAK7392	Aak7392 Human car
c	30	20	100.0	98	13	ADJ08810	Aak708810 Human car
c	31	20	100.0	102	3	AAK13267	Aak13267 Human sec
c	32	20	100.0	102	4	AAK36278	Aak36278 Human car
c	33	20	100.0	102	10	AAK46972	Aak46972 Human car
c	34	20	100.0	102	13	ADJ08390	Adj08390 Human car
c	35	20	100.0	103	4	AAK28603	AAK28603 Genomic s
c	36	20	100.0	103	10	ADG41799	Adg41799 Human res
c	37	20	100.0	103	11	ADI97573	Adi97573 Human res
c	38	20	100.0	104	4	AAK07415	Aak107415 Human rep
c	39	20	100.0	104	5	AAK40769	AAK40769 DNA encod
c	40	20	100.0	104	11	ADJ09975	Adj09975 Human pro
c	41	20	100.0	106	12	ADOS7414	Ado57414 DNA encod
c	42	20	100.0	111	4	AAK74404	Aak74404 Human imm
c	43	20	100.0	113	5	ABAL3542	Abal3542 Human ner
c	44	20	100.0	114	3	AAK21248	Aak21248 Human sec
c	45	20	100.0	114	4	AAK77447	Aak77447 Human imm

ALIGNMENTS

RESULT 1

AEB28959
ID AEB28959 standard; DNA; 20 BP.

AC AEB28959;

XX 22-SEP-2005 (first entry)

DT Human MCOLN1 gene PCR primer PRI F1 SEQ ID NO:3.

DE Human MCOLN1 gene PCR primer PRI F1 SEQ ID NO:3.
KW mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; PCR;
KW primer; ss.

XX Homo sapiens.

OS US2005153300-A1.

PN 14-JUL-2005.

PD 09-JAN-2004; 2004US-00754446.

PF 09-JAN-2004; 2004US-00754446.

PR (QUES-) QUEST DIAGNOSTICS INC.

XX Sun W, Hantash F;

XX WPI; 2005-521160/53.

XX Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by
amplifying the nucleic acid, detecting amplified product with labeled
oligonucleotide probes via a change in fluorescence which indicates the
presence of an ML IV mutant.

XX Claim 1; SEQ ID NO 3; 15pp; English.

XX The invention relates to a method (M1) for detecting the presence of
mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1)
involves contacting the nucleic acid with oligonucleotide primers and
probes, conducting amplification by temperature cycling and monitoring
the accumulation of amplified nucleic acid by detecting an increase in
donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
CC described: (1) an oligonucleotide (ON1) comprising a sequence
CC complementary to the coding or non-coding strand of the mucolin-1
CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
CC sequences, comprising ON1. (M1) is useful for determining the presence of
CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
CC is useful for genetic counseling of individuals at risk for ML IV or at
CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
CC on chromosome 19p13. The present sequence represents a PCR primer for the
CC MCOLN1 gene, which is used in the exemplification of the present
CC invention.

XX
SQ Sequence 20 BP; 1 A; 7 C; 5 G; 7 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
|||||
Db 1 CTTGCTCTGTGGCCAGGCT 20

RESULT 2
ABV75668/c
ID ABV75668 standard; DNA; 24 BP.
XX
AC ABV75668;
XX
DT 28-JAN-2003 (first entry)
XX
DE Human FD16.06 PCR primer 2.
XX
KW Human; FD16.06; cancer; HIV; PCR; primer; ss.
XX
OS Homo sapiens.
XX
PN CN1351034-A.
XX
PD 29-MAY-2002.
XX
PF 26-OCT-2000; 2000CN-00125785.
XX
PR 26-OCT-2000; 2000CN-00125785.
XX
PA (BODE-) BODE GENE DEV CO LTD SHANGHAI.
XX
PI Mao Y, Xie Y;
XX
WPI; 2002-619857/67.
XX
PT Polypeptide-human FD16.06 and polynucleotide for coding it, useful for
PT treating diseases such as cancer and HIV infection.
XX
PS Example 2; Page 16 (Disclosure); 33pp; Chinese.
XX
CC The invention relates to a novel polypeptide, human FD16.06, and the
CC polynucleotide encoding it. The antagonist against the polypeptide is
CC also disclosed. The polypeptide is useful for treating diseases such as
CC cancer and HIV infection. The present sequence represents a PCR primer
CC used to amplify the human FD16.06 gene of the invention
XX
SQ Sequence 24 BP; 9 A; 7 C; 7 G; 1 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
|||||
Db 22 CTTGCTCTGTGGCCAGGCT 3

RESULT 3
AAH48082
ID AAH48082 standard; DNA; 41 BP.
XX
AC AAH48082;
XX
DT 19-SEP-2001 (first entry)
XX
DE Ras GTPase-activating protein 12 probe #2.
XX
KW Ras GTPase-activating protein 12; cytosolic; Antiinflammatory;
KW Immunomodulatory; Anti-HIV; malignant tumour; haemopathy; HIV infection;
KW immunological disease; inflammation; gene therapy; probe; ss.
XX
OS Unidentified.
XX
PN WO200147998-A1.
XX
PD 05-JUL-2001.
XX
PF 25-DEC-2000; 2000WO-CN000680.
XX
PR 27-DEC-1999; 99CN-00125367.
XX
PA (SHAN-) SHANGHAI BIOWINDOW GENE DEV INC.
XX
PI Mao Y, Xie Y;
XX
WPI; 2001-418231/44.
XX
PT RAS GTPase-activating protein 12 and encoded polynucleotide, applicable
PT in diagnosis and treatment of malignant tumor, hemopathy, HIV infection,
PT immunological diseases and various inflammation.
XX
PS Example 7; Page 20; 38pp; Chinese.
XX
CC The present invention relates to Ras GTPase-activating protein 12 and its
CC coding sequence (see AAH48076 and AAG64227). The GTPase-activating
CC protein and its coding sequence are useful in the diagnosis and treatment
CC of malignant tumor, haemopathy, HIV infection, immunological diseases
CC and various inflammations. The present sequence is a probe, which was
CC used in an example from the present invention
XX
SQ Sequence 41 BP; 6 A; 10 C; 12 G; 13 T; 0 U; 0 Other;

Query Match 100.0%; Score 20; DB 4; Length 41;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGGCCAGGCT 20
|||||
Db 13 CTTGCTCTGTGGCCAGGCT 32

RESULT 4
AAS15951/c
ID AAS15951 standard; DNA; 41 BP.
XX
AC AAS15951;
XX
DT 14-FEB-2002 (first entry)
XX
DE Human proteolytic enzyme regulatory protein 11, probe #2.

XX
KW Human; proteolytic enzyme regulatory protein 11; cytosolic; virucidal;
KW immunomodulatory; antiinflammatory; haemostatic; cancer; haemopathy;
KW human immunodeficiency virus; HIV; immunological disease; inflammation;
KW embryonic development disorder; developmental disorder; probe; ss.
XX
OS Homo sapiens.
XX
PN WO200174867-A1.
XX

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 16:41:33 ; Search time 483.04 Seconds
(without alignments)
342.389 Million cell updates/sec

Title: US-10-754-446-3

Perfect score: 20

Sequence: 1 CTTGCTCTGTGCCAGGCT 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 9793542 seqs, 4134689005 residues

Total number of hits satisfying chosen parameters: 19587084

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA Main.*
1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq.*
2: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq.*
3: /cgn2_6/ptodata/1/pubpna/US09A_PUBCOMB.seq.*
4: /cgn2_6/ptodata/1/pubpna/US09B_PUBCOMB.seq.*
5: /cgn2_6/ptodata/1/pubpna/US10A_PUBCOMB.seq.*
6: /cgn2_6/ptodata/1/pubpna/US10B_PUBCOMB.seq.*
7: /cgn2_6/ptodata/1/pubpna/US10C_PUBCOMB.seq.*
8: /cgn2_6/ptodata/1/pubpna/US10D_PUBCOMB.seq.*
9: /cgn2_6/ptodata/1/pubpna/US10E_PUBCOMB.seq.*
10: /cgn2_6/ptodata/1/pubpna/US11_PUBCOMB.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	9	US-10-754-446-3
2	20	100.0	60	8	US-10-804-859-55
3	20	100.0	71	9	US-10-467-851-277
4	20	100.0	72	9	US-10-708-204-6674
5	20	100.0	77	9	US-10-467-851-853
6	20	100.0	78	9	US-10-708-204-6268
7	20	100.0	78	9	US-10-708-204-6998
8	20	100.0	83	9	US-10-708-204-6598
9	20	100.0	84	3	US-09-764-860-972
10	20	100.0	84	5	US-10-074-095-972
11	20	100.0	84	6	US-10-212-872-972
12	20	100.0	84	9	US-10-708-204-6304
13	20	100.0	86	9	US-10-708-204-5760
14	20	100.0	87	9	US-10-708-204-6688
15	20	100.0	88	9	US-10-708-204-6366
16	20	100.0	88	9	US-10-708-204-6626
17	20	100.0	88	9	US-10-708-204-6671
18	20	100.0	88	9	US-10-708-204-6676
19	20	100.0	88	9	US-10-708-204-6689
20	20	100.0	88	9	US-10-708-204-6699
21	20	100.0	88	9	US-10-708-204-6784
22	20	100.0	90	9	US-10-708-204-6710
23	20	100.0	92	9	US-10-708-204-6788

Sequence 6492, Ap
Sequence 6445, Ap
Sequence 6753, Ap
Sequence 6848, Ap
Sequence 5730, Ap
Sequence 6816, Ap
Sequence 6419, Ap
Sequence 6198, Ap
Sequence 2198, Ap
Sequence 2198, Ap
Sequence 6436, Ap
Sequence 5772, Ap
Sequence 6371, Ap
Sequence 6374, Ap
Sequence 6374, Ap
Sequence 7165, Ap
Sequence 1778, Ap
Sequence 1778, Ap
Sequence 1778, Ap
Sequence 1037, Ap
Sequence 1037, Ap
Sequence 1037, Ap

93 9 US-10-708-204-6492
94 9 US-10-708-204-6445
95 9 US-10-708-204-6753
96 9 US-10-708-204-6848
97 9 US-10-708-204-5730
98 9 US-10-708-204-6816
99 9 US-10-708-204-6419
98 3 US-09-764-869-2198
98 5 US-10-091-504-2198
98 6 US-10-227-577-2198
98 9 US-10-708-204-6436
99 9 US-10-708-204-5772
99 9 US-10-708-204-6371
99 9 US-10-708-204-6374
99 9 US-10-708-204-6437
101 9 US-10-708-204-7165
102 3 US-09-764-869-1778
102 5 US-10-091-504-1778
102 6 US-10-227-577-1778
103 3 US-09-764-860-1037
103 5 US-10-074-095-1037
103 6 US-10-212-872-1037

ALIGNMENTS

RESULT 1
US-10-754-446-3
; Sequence 3, Application US/10754446
; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: HANTASH, PERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; TITLE OF INVENTION: MUCOLIPIDOSIS IV MUTATIONS
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: Patent In Ver. 3.2
; SEQ ID NO 3
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-10-754-446-3

Query Match 100.0%; Score 20; DB 9; Length 20;
Best Local Similarity 100.0%; Pred. No. 6.1;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTGCTCTGTGCCAGGCT 20
| | | | | | | | | | | | | | | | | | | | | |
Db 1 CTTGCTCTGTGCCAGGCT 20

RESULT 2
US-10-804-859-55
; Sequence 55, Application US/10804859
; Publication No. US20040265865A1
; GENERAL INFORMATION:
; APPLICANT: The University of Queensland
; TITLE OF INVENTION: A method for identifying effector molecules for gene network inte
; FILE REFERENCE: 2563972/EJH
; CURRENT APPLICATION NUMBER: US/10804,859
; CURRENT FILING DATE: 2004-03-19
; PRIOR APPLICATION NUMBER: US 60/324127
; PRIOR FILING DATE: 2001-09-19
; NUMBER OF SEQ ID NOS: 121
; SOFTWARE: Patent In version 3.0
; SEQ ID NO 55

; LENGTH: 60
; TYPE: DNA
; ORGANISM: human
US-10-804-859-55

Query Match 100.0%; Score 20; DB 8; Length 60;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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DB 7 CTTGCTCTGTGCCAGGCT 26

RESULT 3

US-10-467-851-277
; Sequence 277, Application US/10467851
; Publication No. US20050221303A1
; GENERAL INFORMATION:
; APPLICANT: TELERMAN, Adam
; APPLICANT: AMSON, Robert
; APPLICANT: TUIJNDER, Marcel,
; APPLICANT: SUSINI, Laurent
; TITLE OF INVENTION: SEQUENCES INVOLVED IN PHENOMENA OF TUMOUR SUPPRESSION,
; TITLE OF INVENTION: TUMOUR REVERSION, APOPTOSIS AND/OR VIRUS RESISTANCE
; TITLE OF INVENTION: AND THEIR USE AS MEDICINES
; FILE REFERENCE: 11416-014-999
; CURRENT APPLICATION NUMBER: US/10/467,851
; CURRENT FILING DATE: 2003-08-13
; PRIOR APPLICATION NUMBER: PCT/FR 02/00 543
; PRIOR FILING DATE: 2002-02-13
; PRIOR APPLICATION NUMBER: FR 01/01 925
; PRIOR FILING DATE: 2001-02-13
; NUMBER OF SEQ ID NOS: 1020
; SOFTWARE: Patentin version 3.0
; SEQ ID NO 277
; LENGTH: 71
; TYPE: DNA
; ORGANISM: HOMO SAPIENS
US-10-467-851-277

Query Match 100.0%; Score 20; DB 9; Length 71;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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DB 11 CTTGCTCTGTGCCAGGCT 30

RESULT 4

US-10-708-204-6674/c
; Sequence 6674, Application US/10708204
; Publication No. US20050222399A1
; GENERAL INFORMATION:
; APPLICANT: ROSETTA GENOMICS LTD
; TITLE OF INVENTION: BIOINFORMATICALLY DETECTABLE GROUP OF NOVEL REGULATORY
; TITLE OF INVENTION: OLIGONUCLEOTIDES ASSOCIATED WITH ALZHEIMER'S DISEASE AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 55033
; CURRENT APPLICATION NUMBER: US/10/708,204
; CURRENT FILING DATE: 2004-02-16
; NUMBER OF SEQ ID NOS: 7351
; SOFTWARE: Patentin version 3.2
; SEQ ID NO 6674
; LENGTH: 72
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-10-708-204-6674

Query Match 100.0%; Score 20; DB 9; Length 72;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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DB 67 CTTGCTCTGTGCCAGGCT 48

RESULT 5

US-10-467-851-853/c
; Sequence 853, Application US/10467851
; Publication No. US20050221303A1
; GENERAL INFORMATION:
; APPLICANT: TELERMAN, Adam
; APPLICANT: AMSON, Robert
; APPLICANT: TUIJNDER, Marcel,
; APPLICANT: SUSINI, Laurent
; TITLE OF INVENTION: SEQUENCES INVOLVED IN PHENOMENA OF TUMOUR SUPPRESSION,
; TITLE OF INVENTION: TUMOUR REVERSION, APOPTOSIS AND/OR VIRUS RESISTANCE
; TITLE OF INVENTION: AND THEIR USE AS MEDICINES
; FILE REFERENCE: 11416-014-999
; CURRENT APPLICATION NUMBER: US/10/467,851
; CURRENT FILING DATE: 2003-08-13
; PRIOR APPLICATION NUMBER: PCT/FR 02/00 543
; PRIOR FILING DATE: 2002-02-13
; PRIOR APPLICATION NUMBER: FR 01/01 925
; PRIOR FILING DATE: 2001-02-13
; NUMBER OF SEQ ID NOS: 1020
; SOFTWARE: Patentin version 3.0
; SEQ ID NO 853
; LENGTH: 77
; TYPE: DNA
; ORGANISM: HOMO SAPIENS
US-10-467-851-853

Query Match 100.0%; Score 20; DB 9; Length 77;
Best Local Similarity 100.0%; Pred. No. 5.9;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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DB 39 CTTGCTCTGTGCCAGGCT 20

RESULT 6

US-10-708-204-6268/c
; Sequence 6268, Application US/10708204
; Publication No. US20050222399A1
; GENERAL INFORMATION:
; APPLICANT: ROSETTA GENOMICS LTD
; TITLE OF INVENTION: BIOINFORMATICALLY DETECTABLE GROUP OF NOVEL REGULATORY
; TITLE OF INVENTION: OLIGONUCLEOTIDES ASSOCIATED WITH ALZHEIMER'S DISEASE AND USES
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: 55033
; CURRENT APPLICATION NUMBER: US/10/708,204
; CURRENT FILING DATE: 2004-02-16
; NUMBER OF SEQ ID NOS: 7351
; SOFTWARE: Patentin version 3.2
; SEQ ID NO 6268
; LENGTH: 78
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-10-708-204-6268

Query Match 100.0%; Score 20; DB 9; Length 78;
Best Local Similarity 100.0%; Pred. No. 5.9;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
|||||
DB 78 CTTGCTCTGTGCCAGGCT 59

RESULT 7

US-10-708-204-6998/c

Result No.	Score	Query Match	Length	DB	ID	Description	
1	20	100.0	21	6	US-10-310-914A-206955	Sequence 206955,	
2	20	100.0	21	6	US-10-310-914A-391931	Sequence 391931,	
3	20	100.0	21	6	US-10-310-914A-437321	Sequence 437321,	
4	20	100.0	21	6	US-10-310-914A-578793	Sequence 578793,	
5	20	100.0	21	6	US-10-310-914A-611412	Sequence 611412,	
6	20	100.0	21	6	US-10-310-914A-613165	Sequence 613165,	
7	20	100.0	21	6	US-10-310-914A-762251	Sequence 762251,	
8	20	100.0	21	6	US-10-310-914A-1015245	Sequence 1015245,	
9	20	100.0	21	6	US-10-310-914A-1304732	Sequence 1304732,	
10	20	100.0	22	6	US-10-310-914A-1312897	Sequence 1312897,	
11	20	100.0	22	6	US-10-310-914A-156812	Sequence 156812,	
12	20	100.0	22	6	US-10-310-914A-157391	Sequence 157391,	
13	20	100.0	22	6	US-10-310-914A-184260	Sequence 184260,	
14	20	100.0	22	6	US-10-310-914A-201989	Sequence 201989,	
15	20	100.0	22	6	US-10-310-914A-228639	Sequence 228639,	
16	20	100.0	22	6	US-10-310-914A-532105	Sequence 532105,	
17	20	100.0	22	6	US-10-310-914A-611495	Sequence 611495,	
18	20	100.0	22	6	US-10-310-914A-613215	Sequence 613215,	
19	20	100.0	22	6	US-10-310-914A-732734	Sequence 732734,	
20	20	100.0	22	6	US-10-310-914A-1189956	Sequence 1189956,	
21	20	100.0	22	6	US-10-310-914A-1189957	Sequence 1189957,	
22	20	100.0	22	6	US-10-310-914A-1304781	Sequence 1304781,	
23	20	100.0	22	6	US-10-310-914A-1304782	Sequence 1304782,	

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RESULT 2
US/30-310-914A-391931
; Sequence 391931, Application US/10310914A
; Publication No. US2006003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiler, Kvuizat
; TITLE OF INVENTION: Bioinformatically deter
; TITLE OF INVENTION: uses thereof
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 391931
; LENGTH: 21
; TYPE: RNA

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Result No.	Score	Query %		Length	DB	ID	Description
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C 1	20	100.0	98	3	US-09-513-999C-20008		Sequence 20008, A
C 2	20	100.0	102	3	US-09-513-999C-17342		Sequence 17342, A
C 3	20	100.0	114	3	US-09-513-999C-25323		Sequence 25323, A
C 4	20	100.0	117	3	US-09-513-999C-17724		Sequence 17724, A
C 5	20	100.0	117	3	US-09-513-999C-24469		Sequence 24469, A
C 6	20	100.0	128	3	US-09-513-999C-15044		Sequence 15044, A
C 7	20	100.0	145	3	US-09-621-976-14390		Sequence 14390, A
C 8	20	100.0	145	3	US-09-513-999C-16178		Sequence 16178, A
C 9	20	100.0	145	3	US-09-513-999C-25910		Sequence 25910, A
C 10	20	100.0	151	3	US-09-513-999C-24127		Sequence 24127, A
C 11	20	100.0	152	3	US-09-621-976-11650		Sequence 11650, A
C 12	20	100.0	152	3	US-09-621-976-13583		Sequence 13583, A
C 13	20	100.0	155	3	US-09-513-999C-29581		Sequence 29581, A
C 14	20	100.0	158	3	US-09-621-976-11688		Sequence 11688, A
C 15	20	100.0	158	3	US-09-621-976-12407		Sequence 12407, A
C 16	20	100.0	166	3	US-09-621-976-11848		Sequence 11848, A
C 17	20	100.0	168	3	US-09-621-976-11602		Sequence 11602, A
C 18	20	100.0	170	3	US-09-621-976-11644		Sequence 11644, A
C 19	20	100.0	170	3	US-09-621-976-11777		Sequence 11777, A
C 20	20	100.0	170	3	US-09-621-976-11869		Sequence 11869, A
C 21	20	100.0	170	3	US-09-621-976-12011		Sequence 12011, A
C 22	20	100.0	170	3	US-09-621-976-12098		Sequence 12098, A
C 23	20	100.0	170	3	US-09-621-976-12260		Sequence 12260, A
C 24	20	100.0	170	3	US-09-621-976-12260		Sequence 12260, A

; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 17342
; LENGTH: 102
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-17342

Query Match 100.0%; Score 20; DB 3; Length 102;
Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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Db 78 CTTGCTCTGTGCCAGGCT 97

RESULT 3

US-09-513-999C-25323/c
; Sequence 25323, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 25323
; LENGTH: 114
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-25323

Query Match 100.0%; Score 20; DB 3; Length 114;
Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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Db 91 CTTGCTCTGTGCCAGGCT 72

RESULT 4

US-09-513-999C-17724
; Sequence 17724, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 17724
; LENGTH: 117
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 9

; OTHER INFORMATION: k=g or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 11
; OTHER INFORMATION: k=g or t
US-09-513-999C-17724

Query Match 100.0%; Score 20; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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Db 25 CTTGCTCTGTGCCAGGCT 44

RESULT 5

US-09-513-999C-24469/c
; Sequence 24469, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 24469
; LENGTH: 117
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-24469

Query Match 100.0%; Score 20; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 3.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTGCTCTGTGCCAGGCT 20
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RESULT 6

US-09-513-999C-15044
; Sequence 15044, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
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; LENGTH: 128
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-15044

Query Match 100.0%; Score 20; DB 3; Length 128;
Best Local Similarity 100.0%; Pred. No. 3.7;

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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Maximum Match 100%

Listing first 45 summaries

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4: gb_on.*
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6: gb_pat.*
7: gb_ph.*
8: gb_pr.*
9: gb_ro.*
10: gb_sts.*
11: gb_sy.*
12: gb_un.*
13: gb_vi.*
14: gb_htg.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	22	100.0	1740	6 AX280021	AX280021 Sequence
C 3	22	100.0	1829	6 CO719994	CO719994 Sequence
C 4	22	100.0	2004	8 AF249319	AF249319 Homo sapi
C 5	22	100.0	2037	8 HSA233970	HSA233970 Homo sapi
C 6	22	100.0	2049	8 AK026102	AK026102 Homo sapi
C 7	22	100.0	2051	8 AF287269	AF287269 Homo sapi
C 8	22	100.0	2052	8 AK083508	AK083508 Sequence
C 9	22	100.0	2063	8 AK222673	AK222673 Homo sapi
C 10	22	100.0	2078	8 BC005149	BC005149 Homo sapi
C 11	22	100.0	2094	6 BD233734	BD233734 31 human
C 12	22	100.0	2095	6 AX280019	AX280019 Sequence
C 13	22	100.0	2272	8 HSA293659	HSA293659 Homo sapi
C 14	22	100.0	13270	8 AF287270	AF287270 Homo sapi
C 15	22	100.0	155645	14 AC021153	AC021153 Homo sapi
C 16	22	100.0	173126	8 AC008878	AC008878 Homo sapi
C 17	20.4	92.7	2065	8 AB125179	AB125179 Macaca fa
C 18	20	90.9	110000	1 AB014295_10	Continuation (11 o

C 19	20	90.9	349980	6 AX492785	AX492785 Sequence
C 20	20	90.9	349980	6 AX553952	AX553952 Sequence
C 21	17.8	80.9	717	10 BV552653	BV552653 S221P6043
C 22	17.8	80.9	734	10 BV539631	BV539631 G591P6424
C 23	17.8	80.9	767	10 BV534108	BV534108 G591P6224
C 24	17.8	80.9	775	10 BV539839	BV539839 G591P6408
C 25	17.8	80.9	826	10 BV532664	BV532664 G591P6209
C 26	17.8	80.9	1347	13 DQ103713	DQ103713 Influenza
C 27	17.8	80.9	3666	1 AB050784	AB050784 Spingomo
C 28	17.8	80.9	7722	1 AB032203	AB032203 Spingomo
C 29	17.8	80.9	24756	14 AC010403	AC010403 Homo sapi
C 30	17.8	80.9	69678	8 AC091573	AC091573 Homo sapi
C 31	17.8	80.9	143803	5 BX470131	BX470131 Zebrafish
C 32	17.8	80.9	157076	8 AC019031	AC019031 Homo sapi
C 33	17.8	80.9	174578	8 AC013287	AC013287 Homo sapi
C 34	17.8	80.9	177301	8 AC078868	AC078868 Homo sapi
C 35	17.8	80.9	182182	5 BX890617	BX890617 Zebrafish
C 36	17.8	80.9	212671	14 AC084849	AC084849 Homo sapi
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C 41	17.4	79.1	4483	1 STYHISTO	J01805 S. typhimur
C 42	17.4	79.1	8516	1 D88362	D88362 Arthrobacte
C 43	17.4	79.1	20518	1 AE008805	AE008805 Salmonell
C 44	17.4	79.1	40699	2 CEZK637	Z1115 Caenorhabdi
C 45	17.4	79.1	82568	15 AP004906	AP004906 Lotus cor

ALIGNMENTS

RESULT 1
AF305572S3/c AF305572S3 790 bp DNA linear PRI 26-DEC-2000
LOCUS Homo sapiens mucolin 1 (MCOLN1) gene, exons 3 and 4.
DEFINITION Homo sapiens mucolin 1 (MCOLN1) gene, exons 3 and 4.
ACCESSION AF305574
VERSION AF305574.1 GI:11991200
KEYWORDS
SEGMENT
SOURCE 3 of 8
ORGANISM Homo sapiens (human)

REFERENCE 1 (bases 1 to 790)
AUTHORS Bargal,R., Avidan,N., Ben-Asher,E., Olender,Z., Zeigler,M., Frumkin,A., Raas-Rothschild,A., Glusman,G., Lancet,D. and Bach,G.
TITLE Identification of the gene causing mucopolidiosis type IV
JOURNAL Nat. Genet. 26 (1), 118-123 (2000)
PUBMED 10973263
REFERENCE 2 (bases 1 to 790)
AUTHORS Bargal,R., Avidan,N., Ben-Asher,E., Olender,A., Zeigler,M., Frumkin,A., Raas-Rothschild,A., Glusman,G., Lancet,D. and Bach,G.
TITLE Direct Submission
JOURNAL Submitted (13-SEP-2000) Molecular Genetics, The Weizmann Institute of Science, P. O. Box 26, Rehovot 76100, Israel
FEATURES
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Query Match 100.0%; Score 22; DB 8; Length 790;
Best Local Similarity 100.0%; Pred. No. 0.87; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
Db 750 TAACCACCATCGGATCAATGTC 729

RESULT 2
LOCUS AX280021/c 1740 bp DNA linear PAT 02-NOV-2001
DEFINITION Sequence 3 from Patent WO0177331.
ACCESSION AX280021
VERSION AX280021.1 GI:16607475
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE 1
AUTHORS Curtis, R.A. and Silos-Santiago, I.
TITLE Human tripartite calcium channel protein-2 (tlcc-2)
JOURNAL Patent: WO 017331-A 3 18-OCT-2001;
MILLENNIUM PHARMACEUTICALS, INC. (US)

FEATURES
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/codon_start=1
/protein_id="CAD10547.1"
/db_xref="GI:16607476"
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HLFLGYSQADDTFAAYTREQLYQAIHFAVDQYLALPDVSLGRYAYVGGGDPWTNG
SGLALCQRYHGHVDPANDTFIDPMVVTDCIQDPPPPPPSDLLTLLSSSY
KNLTKFHLVNTIHFRLKTNLQSLINNEIPDCYTFVSLITFDNKAHSGRIPI SLE
TQAHIOECKHPSPVQHGDSFRLLFDVWVILTCSLFLLCARSLRGLLQNEFVGFM
WRQGRVLSLWERLEFVNGWYLLVTSVLTSGTIMGIEAKNLASYDVCISLLGT
STLVWGVVIRYLTFFHNHNLIALTLRALVPSVNRFCVAVIYLGVCFCGWLVLGY
HVFRLSLWSECLFSLNGDDMPVTFPAAQOQGRSSLVWFLFSQLYLSFISLFYIM
VLSLFTALITGAYDTIKHPGGAEESELOQYIAQCQDSPTSGKFRGSGSACSLCC
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CDS
1. .1740
/notes="unnamed protein product"
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LKYPFSPCKFRAGKPKMLQVVKILVTVQLFGLSNQLAVTFREENTIAFR
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SGLALCQRYHGHVDPANDTFIDPMVVTDCIQDPPPPPPSDLLTLLSSSY
KNLTKFHLVNTIHFRLKTNLQSLINNEIPDCYTFVSLITFDNKAHSGRIPI SLE
TQAHIOECKHPSPVQHGDSFRLLFDVWVILTCSLFLLCARSLRGLLQNEFVGFM
WRQGRVLSLWERLEFVNGWYLLVTSVLTSGTIMGIEAKNLASYDVCISLLGT
STLVWGVVIRYLTFFHNHNLIALTLRALVPSVNRFCVAVIYLGVCFCGWLVLGY
HVFRLSLWSECLFSLNGDDMPVTFPAAQOQGRSSLVWFLFSQLYLSFISLFYIM
VLSLFTALITGAYDTIKHPGGAEESELOQYIAQCQDSPTSGKFRGSGSACSLCC
CGRDPSEHSLVN"

ORIGIN
Query Match 100.0%; Score 22; DB 6; Length 1740;
Best Local Similarity 100.0%; Pred. No. 0.96;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
Db 568 TAACCACCATCGGATCAATGTC 547

RESULT 3
LOCUS CQ719994/c 1829 bp DNA linear PAT 03-FEB-2004
DEFINITION Sequence 5928 from Patent WO02068579.
ACCESSION CQ719994
VERSION CQ719994.1 GI:42280851
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.

REFERENCE 1
AUTHORS Venter, C.J., Adams, M.C., Li, P.W. and Myers, E.W.
TITLE Kits, such as nucleic acid arrays, comprising a majority of
human exons or transcripts, for detecting expression and other uses
thereof
JOURNAL Patent: WO 02068579-A 5928 06-SEP-2002;

PE Corporation (NY) (US)
Location/Qualifiers
1. .1829
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 0.96;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
Db 537 TAACCACCATCGGATCAATGTC 516

RESULT 4
LOCUS AF249319/c 2004 bp mRNA linear PRI 22-SEP-2000
DEFINITION Homo sapiens mucopolipidosis type IV protein (MLIV) mRNA, complete
cds.
ACCESSION AF249319
VERSION AF249319.1 GI:9971787
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
1 (bases 1 to 2004)
Bargal, R., Avidan, N., Ben-Asher, E., Olender, Z., Zeigler, M.,
Frumkin, A., Rases-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
IDENTIFICATION of the gene causing mucopolipidosis type IV
NAT. GENET. 26 (1), 118-123 (2000)
10973263
2 (bases 1 to 2004)
Bargal, R., Avidan, N., Ben-Asher, E., Olender, Z., Zeigler, M.,
Frumkin, A., Rases-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
DIRECT SUBMISSION Molecular Genetics, The Weizmann Institute
Submitted (26-MAR-2000) of Science, P. O. Box 26, Rehovot 76100, Israel
Location/Qualifiers
1. .2004
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
1. .2004
/genes="MLIV"
85. .1827
/gene="MLIV"
/note="causes mucopolipidosis type IV"
/codon_start=1
/product="mucopolipidosis type IV protein"
/protein_id="AAG10422.1"
/db_xref="GI:9971788"

ORIGIN
Query Match 100.0%; Score 22; DB 8; Length 2004;
Best Local Similarity 100.0%; Pred. No. 0.97;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 2460.3 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-2

Perfect score: 22

Sequence: 1 taaccaccatcgatcaatgctc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST.*

1: gb_est1.*
2: gb_est2.*
3: gb_est3.*
4: gb_hic.*
5: gb_est4.*
6: gb_est5.*
7: gb_est6.*
8: gb_est7.*
9: gb_gss1.*
10: gb_gss2.*
11: gb_gss3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	22	100.0	423	1	AW969529
C 2	22	100.0	425	3	BP391004
C 3	22	100.0	439	8	T66341
C 4	22	100.0	480	1	AW328429
C 5	22	100.0	510	2	BG272754
C 6	22	100.0	554	3	BP265945
C 7	22	100.0	580	3	BP266237
C 8	22	100.0	581	3	BP332355
C 9	22	100.0	657	3	B1819028
C 10	22	100.0	660	3	B1836168
C 11	22	100.0	660	3	B1907877
C 12	22	100.0	663	3	B1523736
C 13	22	100.0	663	3	B1836493
C 14	22	100.0	666	3	BM005772
C 15	22	100.0	666	6	CA391526
C 16	22	100.0	672	3	B1490022
C 17	22	100.0	678	8	DR001457
C 18	22	100.0	683	6	CF132757
C 19	22	100.0	696	2	BF529860
C 20	22	100.0	697	3	BM049487
C 21	22	100.0	705	3	B1822028
C 22	22	100.0	709	3	B1825945

C 23	22	100.0	743	3	BI771045
C 24	22	100.0	790	6	CB988285
C 25	22	100.0	810	3	BI753742
C 26	22	100.0	818	2	BG831705
C 27	22	100.0	846	5	BX383671
C 28	22	100.0	873	6	CD558014
C 29	22	100.0	890	5	BX399044
C 30	22	100.0	901	5	BUS42887
C 31	22	100.0	902	5	BQ901107
C 32	22	100.0	909	5	BUS43879
C 33	22	100.0	910	5	BUS52585
C 34	22	100.0	962	5	BX402300
C 35	22	100.0	998	3	BI824195
C 36	22	100.0	998	3	BM905707
C 37	22	100.0	1026	5	BX364018
C 38	22	100.0	1047	2	BE795087
C 39	22	100.0	1050	5	BX462609
C 40	22	100.0	1078	5	BX362786
C 41	22	100.0	1144	5	BX386808
C 42	22	100.0	1743	10	AY410283
C 43	22	100.0	1996	4	CR622331
C 44	20.4	92.7	951	5	BU195645
C 45	20	90.9	540	3	BI516259

ALIGNMENTS

RESULT 1
AW969529/c
LOCUS AW969529 423 bp mRNA linear EST 01-JUN-2000
DEFINITION EST381606 MAGE resequences, MAGK Homo sapiens cDNA, mRNA sequence.
ACCESSION AW969529
VERSION AW969529.1 GI:8159373
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 423)
AUTHORS Hegde,P., Qi,R., Abernathy,K., Dharap,S., Gaspard,R., Gay,C., Holt,I.E., Saeed,A.I., Sharov,V., Lee,N.H., Yeatman,T.J. and Quackenbush,J.
TITLE Assessment of gene expression patterns in a model of colon tumor metastasis using a 19,200 element cDNA microarray
JOURNAL Unpublished (2000)
COMMENT Contact: John Quackenbush
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 3528
Fax: 301 838 0208
Email: johnq@tigr.org
Plate: 286
Seq primer: Forward.
Location/Qualifiers
1 . 423
/organism="Homo sapiens"
/mol_type="mRNA"
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/clone_lib="MAGE resequences, MAGK"
/note="Vector: pBluescriptSKm"

Query Match 100.0%; Score 22; DB 1; Length 423;
Best Local Similarity 100.0%; Pred. No. 2.;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGATCAATGTC 22
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DB 96 TAACCACCATCGATCAATGTC 75

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RESULT 2
BP391004/c
LOCUS
DEFINITION BP391004 Homo sapiens pancreatic islet Homo sapiens cDNA clone
h-t-232-79 3', mRNA sequence.
ACCESSION BP391004
VERSION BP391004.1 GI:58414498
SOURCE EST.
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE
1 (bases 1 to 425)
Takeda,J., Jin,L. and Horikawa,Y.
TITLE Expression profile of mRNAs from human pancreatic islet tumors
JOURNAL Unpublished (2005)
COMMENT Contact: Yukio Horikawa
Laboratory of Molecular Genetics
Institute for Molecular and Cellular Regulation, Gunma University
Showa-machi 3-39-15, Maebashi, Gunma, 371-8512, Japan
Tel: 81-27-220-8832
Fax: 81-27-220-8889
Email: yhorikawa@showa.gunma-u.ac.jp.
FEATURES
source
1..425
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="h-t-232-79"
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/clone_lib="Homo sapiens pancreatic islet"

ORIGIN
Query Match 100.0%; Score 22; DB 3; Length 425;
Best Local Similarity 100.0%; Pred. No. 2.7;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
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Db 73 TAACCACCATCGGATCAATGTC 52
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RESULT 3
T66341/c
LOCUS
DEFINITION T66341 439 bp mRNA linear EST 07-MAR-1995
yc7806.r1 Soares infant brain INIB Homo sapiens cDNA clone
IMAGE:21977 5', mRNA sequence.
T66341
ACCESSION T66341.1 GI:675386
VERSION EST.
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE
1 (bases 1 to 439)
Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M.,
Holman,M., Hultman,M., Kucaba,T., Le,M., Lemmon,G., Marra,M.,
Parsons,J., Rifkin,L., Rolfing,T., Soares,M., Tan,F.,
Trevaskis,E., Waterston,R., Williamson,A., Wohlmann,P. and
Wilson,R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT Contact: Willson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 2264
High quality sequence stops: 345 Source: IMAGE Consortium, LLNL This
clone is available royalty-free through LLNL ; contact the IMAGE

Consortium (info@image.llnl.gov) for further information.
Insert Length: 2264 Std Error: 0.00
Seq primer: M13RP1
High quality sequence stop: 345.
Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:394324"
/db_xref="taxon:9606"
/clone="IMAGE:21977"
/sex="female"
/dev_stage="73 days post natal"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares infant brain INIB"
/notes="Organ: whole brain; Vector: Lafmid BA; Site 1: Not
I; Site 2: Hind III; 1st strand cDNA was primed with a Not
I - oligo(dT) primer [5'
AACTGGAGAAATTCGGCGCCAGGAAATTTTTTTTTTTTTTTT 3'];
double-stranded cDNA was ligated to Hind III adaptors
(Pharmacia), digested with Not I and directionally cloned
into the Not I and Hind III sites of the Lafmid BA vector.
Library went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."

ORIGIN
Query Match 100.0%; Score 22; DB 8; Length 439;
Best Local Similarity 100.0%; Pred. No. 2.7;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
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Db 309 TAACCACCATCGGATCAATGTC 288
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RESULT 4
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LOCUS
DEFINITION AW328429 480 bp mRNA linear EST 28-JAN-2000
ds02e06.x1 NIH_MGC_4 Homo sapiens cDNA clone IMAGE:2847443 5', mRNA
sequence.
AW328429
ACCESSION AW328429.1 GI:6798925
VERSION EST.
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE
1 (bases 1 to 480)
NIH-MGC http://mgi.nci.nih.gov/.
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE Unpublished (1999)
JOURNAL
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgabbs@email.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Edge Biosystems
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: NIH Intramural Sequencing Center (NISC)
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Plate: LLCM0030 row: I column: 12
Seq primer: -21M13 forward primer (ABI).
Location/Qualifiers
1..480
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2847443"
/cell_line="MGC1"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH MGC 4"
/notes="Organ: Cervix; Vector: pOTB7a; Site 1: Scl;
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 14:15:14 ; Search time 298.496 Seconds
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491.207 Million cell updates/sec

Title: US-10-754-446-2

Perfect score: 22

Sequence: 1 taaccaccatcgatcaatgctc 22

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4996997 seqs, 332346308 residues

Total number of hits satisfying chosen parameters: 9993994

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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1: Geneseqn1980s.*

2: Geneseqn1990s.*

3: Geneseqn2000s.*

4: Geneseqn2001as.*

5: Geneseqn2001bs.*

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7: Geneseqn2002bs.*

8: Geneseqn2003as.*

9: Geneseqn2003bs.*

10: Geneseqn2003cs.*

11: Geneseqn2003ds.*

12: Geneseqn2004as.*

13: Geneseqn2004bs.*

14: Geneseqn2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	22	100.0	22	14	AEB28958 Human MCO
2	22	100.0	1619	4	Aa161022 Human pol
3	22	100.0	1740	6	Aa171700 Human TRP
4	22	100.0	1740	6	ABL40755 Human TLC
5	22	100.0	1741	4	Aa159236 Human pol
6	22	100.0	1743	9	ADB84284 Human muc
7	22	100.0	1743	14	ADV66231 TRP-like
8	22	100.0	2051	13	ADP25150 PRO polyp
9	22	100.0	2052	4	AaF81753 Human mem
10	22	100.0	2092	5	AaF72274 DNA encod
11	22	100.0	2092	6	ABL90358 Human pol
12	22	100.0	2094	3	AaA39067 Human sec
13	22	100.0	2095	6	Aa171699 Human TRP
14	22	100.0	2095	6	ABL40754 Human TLC
15	22	100.0	2095	14	ADV66229 TRP-like
16	22	100.0	2140	12	ADQ24925 Human sof
17	22	100.0	11365	4	AaK73827 Human imm
18	22	100.0	13270	9	ADB84283 Human muc
19	22	100.0	20046	4	AaK73826 Human imm

C 20	20	90.9	349980	6	ABQ81847
C 21	18.4	83.6	2860	5	AAS65933
C 22	17.4	79.1	682	8	ACA49409
C 23	17.4	79.1	687	8	ACA51484
C 24	17.4	79.1	3153	2	AAV19895
C 25	17.2	78.2	963	13	ADT46459
C 26	16.8	76.4	447	13	ADT43994
C 27	16.8	76.4	705	3	AAS50958
C 28	16.8	76.4	2505	13	ADU01691
C 29	16.4	74.5	1153	4	ABL16397
C 30	16.4	74.5	1274	5	AAS79165
C 31	16.4	74.5	3208	4	ABL16396
C 32	16.4	74.5	4106	4	ABL17062
C 33	16.4	74.5	110000	11	ADM27081_07
C 34	16.2	73.6	230	14	ADY99718
C 35	16.2	73.6	531	12	ACH70153
C 36	16.2	73.6	1035	4	AAD08299
C 37	16.2	73.6	1228	11	ADM22430
C 38	16.2	73.6	1339	13	ADT16087
C 39	16.2	73.6	1638	13	ADT46663
C 40	16.2	73.6	2442	4	ABL08173
C 41	16.2	73.6	2609	13	ADX34421
C 42	16.2	73.6	3825	11	ACL28924
C 43	16.2	73.6	4442	4	ABL08172
C 44	16.2	73.6	4816	10	ADE54538
C 45	16.2	73.6	4816	10	ADE54542

ALIGNMENTS

RESULT 1

AEB28958

ID AEB28958 standard; DNA; 22 BP.

AC AEB28958;

XX 22-SEP-2005 (first entry)

XX Human MCOLN1 gene PCR primer IVS-R SEQ ID NO:2.

XX mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; PCR; primer; ss.

XX Homo sapiens.

XX US2005153300-A1.

XX 14-JUL-2005.

XX 09-JAN-2004; 2004US-00754446.

XX 09-JAN-2004; 2004US-00754446.

XX (QUES-) QUEST DIAGNOSTICS INC.

XX Sun W, Hantash F;

XX WPI; 2005-521160/53.

XX Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by amplifying the nucleic acid, detecting amplified product with labeled oligonucleotide probes via a change in fluorescence which indicates the presence of an ML IV mutant.

XX Claim 1; SEQ ID NO 2; 15pp; English.

XX The invention relates to a method (M1) for detecting the presence of mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1) involves contacting the nucleic acid with oligonucleotide primers and probes, conducting amplification by temperature cycling and monitoring the accumulation of amplified nucleic acid by detecting an increase in donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
 CC described: (1) an oligonucleotide (ON1) comprising a sequence
 CC complementary to the coding or non-coding strand of the mucolipin-1
 CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
 CC sequences, comprising ON1. (M1) is useful for determining the presence of
 CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
 CC is useful for genetic counseling of individuals at risk for ML IV or at
 CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
 CC on chromosome 19p13. The present sequence represents a PCR primer for the
 CC MCOLN1 gene, which is used in the exemplification of the present
 CC invention.

XX SQ Sequence 22 BP; 7 A; 7 C; 3 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 14; Length 22;
 Best Local Similarity 100.0%; Pred. No. 0.3;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
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 Db 1 TAACCACCATCGGATCAATGTC 22

RESULT 2
 AAI61022
 ID AAI61022 standard; cDNA; 1619 BP.

XX AAI61022;

DT 22-OCT-2001 (first entry)

XX Human polynucleotide SEQ ID NO 5011.

XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
 KW peripheral nervous system; neuropathy; central nervous system; CNS;
 KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
 KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
 KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
 KW leukaemia; ss.

XX Homo sapiens.

XX WO200153312-A1.

XX 26-JUL-2001.

XX 26-DEC-2000; 2000WO-US034263.

XX 23-DEC-1999; 99US-00471275.

XX 21-JAN-2000; 2000US-00488725.

XX 25-APR-2000; 2000US-00552317.

XX 20-JUN-2000; 2000US-00598042.

XX 19-JUL-2000; 2000US-00620312.

XX 03-AUG-2000; 2000US-00653450.

XX 14-SEP-2000; 2000US-00662191.

XX 19-OCT-2000; 2000US-00693036.

XX 29-NOV-2000; 2000US-00727344.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;

XX Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J, Zhao QA;

XX Zhou P, Goodrich R, Drmanac RT;

XX WPI; 2001-442253/47.

XX P-PSDB; AAM41866.

XX Novel nucleic acids and polypeptides, useful for treating disorders such

XX as central nervous system injuries.

XX Claim 1; SEQ ID NO 5011; 10078pp; English.

XX The invention relates to human nucleic acids (AAI57798-AAI61369) and the

XX

CC

CC encoded polypeptides (AAM38642-AAM42213) with nootropic,
 CC immunosuppressant and cytostatic activity. The polynucleotides are useful
 CC in gene therapy. A composition containing a polypeptide or polynucleotide
 CC of the invention may be used to treat diseases of the peripheral nervous
 CC system, such as peripheral nervous injuries, peripheral neuropathy and
 CC localised neuropathies and central nervous system diseases, such as
 CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
 CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
 CC utilisation of the activities such as: Immune system suppression.
 CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
 CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
 CC assays for receptor activity, arthritis and inflammation, leukaemia and
 CC C.N.S disorders. Note: The sequence data for this patent did not form
 CC part of the printed specification

XX SQ Sequence 1619 BP; 350 A; 450 C; 518 G; 301 T; 0 U; 0 Other;

Query Match 100.0%; Score 22; DB 4; Length 1619;
 Best Local Similarity 100.0%; Pred. No. 0.51;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
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 Db 1382 TAACCACCATCGGATCAATGTC 1403

RESULT 3

AAI71700/c

ID AAI71700 standard; cDNA; 1740 BP.

XX AAI71700;

XX 29-JAN-2002 (first entry)

XX Human TRP-like calcium channel TLCC-2 coding sequence #2.

XX Human; TLCC-2; TRP-like calcium channel; membrane excitability;
 KW nociception; nootropic; neuroprotective; antiparkinsonian; cytostatic;
 KW hypotensive; antidepressant; analgesic; anticonvulsant; tranquiliser;
 KW Parkinson's disease; Huntington's disease; multiple sclerosis;
 KW Gilles de la Tourette's syndrome; autonomic function disorder; cancer;
 KW neuroleptic; gene therapy; Alzheimer's disease; CNS disorder; ss.

XX Homo sapiens.

XX Key Location/Qualifiers
 XX CDS 1..1740

XX /*tag= a
 XX /product= "TLCC-2"
 XX /partial

XX WO200177331-A1.

XX 18-OCT-2001.

XX 06-APR-2001; 2001WO-US011442.

XX 07-APR-2000; 2000US-00544797.

XX (MILL-) MILLENIUM PHARM INC.

XX Curtis RAJ, Silos-Santiago I;

XX WPI; 2002-010913/01.

XX P-PSDB; AAM51858.

XX Novel isolated human transient receptor potential-like calcium channel
 PT protein-2 useful for treating Alzheimer's disease, depression, amnesia,
 PT pain disorder, and cancer.

XX Claim 1; Fig 1; 148pp; English.

XX The present invention relates to the protein and coding sequences of

CC

1	22	100.0	22	9	US-10-754-446-2	Sequence 2, Appli
2	22	100.0	1400	10	US-11-060-756-2850	Sequence 2850, A
3	22	100.0	1400	10	US-11-060-756-7122	Sequence 7122, A
4	22	100.0	1740	3	US-09-828-466-3	Sequence 3, Appli
5	22	100.0	1740	3	US-10-103-458-3	Sequence 3, Appli
6	22	100.0	1740	5	US-10-103-458-3	Sequence 3, Appli
7	22	100.0	1743	8	US-10-782-695-6	Sequence 6, Appli
8	22	100.0	2051	3	US-09-851-494B-2	Sequence 2, Appli
9	22	100.0	2052	3	US-09-965-529-50	Sequence 50, Appli
10	22	100.0	2052	3	US-09-969-680A-50	Sequence 50, Appli
11	22	100.0	2052	10	US-11-048-692-50	Sequence 50, Appli
12	22	100.0	2092	6	US-10-264-237-920	Sequence 920, App
13	22	100.0	2092	9	US-10-450-763-8078	Sequence 8078, Ap
14	22	100.0	2094	3	US-09-820-893-26	Sequence 26, Appli
15	22	100.0	2095	3	US-10-607-563-26	Sequence 26, Appli
16	22	100.0	2095	5	US-09-828-466-1	Sequence 1, Appli
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18	22	100.0	2095	8	US-10-782-695-4	Sequence 4, Appli
19	22	100.0	2140	8	US-10-723-860-7745	Sequence 7745, Ap
20	22	100.0	13270	3	US-09-851-494B-1	Sequence 1, Appli
21	20	90.9	25	10	US-11-060-756-171643	Sequence 171643,
22	20	90.9	2256646	7	US-10-470-565-1	Sequence 1, Appli
23	18.4	83.6	2860	9	US-10-450-763-1737	Sequence 1737, Ap
24	17.4	79.1	682	7	US-10-282-122A-37279	Sequence 37279, A

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; Sequence 2850, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles
; TITLE OF INVENTION: Target Genes
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 2850

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; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-2850

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Best Local Similarity 100.0%; Pred. No. 0.92;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 42 TAACCACCATCGGATCAATGTC 21

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; Sequence 7122, Application US/11060756
; Publication No. US20050221354A1
; GENERAL INFORMATION:
; APPLICANT: Wyeth
; APPLICANT: Mounts, William Martin
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug
; FILE REFERENCE: Target Genes
; FILE REFERENCE: AM101083 (031896-042000)
; CURRENT APPLICATION NUMBER: US/11/060,756
; CURRENT FILING DATE: 2005-02-18
; NUMBER OF SEQ ID NOS: 303284
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 7122
; LENGTH: 1400
; TYPE: DNA
; ORGANISM: Homo sapiens
US-11-060-756-7122

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RESULT 4
US-09-828-466-3/c
; Sequence 3, Application US/09828466
; Patent No. US20020035056A1
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Silos-Santiago, Immaculada
; TITLE OF INVENTION: 54420, A NOVEL HUMAN CALCIUM CHANNEL
; FILE REFERENCE: MNI-125CP
; CURRENT APPLICATION NUMBER: US/09/828,466
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 09/544,797
; PRIOR FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 1740
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1740)
US-09-828-466-3

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Best Local Similarity 100.0%; Pred. No. 0.94;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 568 TAACCACCATCGGATCAATGTC 547

RESULT 5
US-10-103-458-3/c
; Sequence 3, Application US/10103458
; Publication No. US20020197680A1
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; TITLE OF INVENTION: 54420, A NOVEL HUMAN CALCIUM CHANNEL
; FILE REFERENCE: MNI-125
; CURRENT APPLICATION NUMBER: US/10/103,458
; CURRENT FILING DATE: 2002-03-22
; PRIOR APPLICATION NUMBER: US/09/544,797
; PRIOR FILING DATE: PEI0R FILING DATE: 2000-04-07
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
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; TYPE: DNA
; ORGANISM: Homo sapiens
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; NAME/KEY: CDS
; LOCATION: (1)..(1740)
US-10-103-458-3

Query Match      100.0%; Score 22; DB 5; Length 1740;
Best Local Similarity 100.0%; Pred. No. 0.94;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
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Db 568 TAACCACCATCGGATCAATGTC 547

RESULT 6
US-10-782-695-6/c
; Sequence 6, Application US/10782695
; Publication No. US20040248160A1
; GENERAL INFORMATION:
; APPLICANT: Gluckemann, Maria A.
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Tsai, Fong-Ying
; APPLICANT: Hodge, Martin R.
; APPLICANT: Meyers, Rachel E.
; APPLICANT: MacBeth, Kyle J.
; APPLICANT: Bandaru, Rajasekhar
; TITLE OF INVENTION: NOVEL 14275, 54420, 8797, 27439, 68730,
; FILE REFERENCE: MPI04-0020NMIM
; CURRENT APPLICATION NUMBER: US/10/782,695
; CURRENT FILING DATE: 2004-02-19
; PRIOR APPLICATION NUMBER: US 09/945,254
; PRIOR FILING DATE: 2001-08-31
; PRIOR APPLICATION NUMBER: US 60/229,829
; PRIOR FILING DATE: 2000-08-31
; PRIOR APPLICATION NUMBER: US 09/945,301
; PRIOR FILING DATE: 2001-08-31
; PRIOR APPLICATION NUMBER: US 60/229,301
; PRIOR FILING DATE: 2000-09-01
; PRIOR APPLICATION NUMBER: US 10/007,399
; PRIOR FILING DATE: 2001-11-05
; PRIOR APPLICATION NUMBER: US 09/390,039
; PRIOR FILING DATE: 1999-09-03
; PRIOR APPLICATION NUMBER: US 09/146,416
; PRIOR FILING DATE: 1998-09-03
; PRIOR APPLICATION NUMBER: US 10/024,036
; PRIOR FILING DATE: 2001-12-17
; PRIOR APPLICATION NUMBER: US 60/258,222
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: US 10/103,458
; PRIOR FILING DATE: 2002-03-22
; Remaining Prior Application data removed - See File Wrapper or PALM.
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Title: US-10-754-446-2

Perfect score: 22

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 2	16.8	76.4	1107	6	US-10-750-623-53745 Sequence 53745, A
C 3	16.2	73.6	1458	7	US-11-133-360-4 Sequence 4, Appl
C 4	16.2	73.6	1786	6	US-10-750-185-46643 Sequence 46643, A
C 5	16.2	73.6	1786	6	US-10-750-623-46643 Sequence 46643, A
C 6	15.8	71.8	600	7	US-11-128-061-3932 Sequence 3932, Ap
C 7	15.8	71.8	669	7	US-11-128-061-290 Sequence 290, App
C 8	15.8	71.8	2081	6	US-10-750-185-27573 Sequence 27573, A
C 9	15.8	71.8	2081	6	US-10-750-623-27573 Sequence 27573, A
C 10	15.6	70.9	506	7	US-11-128-061-2800 Sequence 2800, Ap
C 11	15.6	70.9	506	7	US-11-128-061-6442 Sequence 6442, Ap
C 12	15.6	70.9	98345	7	US-11-112-908-36 Sequence 36, Appl
C 13	15.6	70.9	127340	7	US-11-112-908-35 Sequence 35, Appl
C 14	15.6	70.9	157224	7	US-11-112-908-51 Sequence 51, Appl
C 15	15.6	70.9	170189	7	US-11-112-908-50 Sequence 50, Appl
C 16	15.4	70.0	88873	6	US-10-995-561-13383 Sequence 13383, A
C 17	15.4	70.0	146656	7	US-11-121-086-68 Sequence 68, Appl
C 18	15.4	70.0	154452	7	US-11-121-086-74 Sequence 74, Appl
C 19	15.4	70.0	190882	7	US-11-121-086-69 Sequence 69, Appl
C 20	15.2	69.1	25	7	US-11-136-527-311442 Sequence 311442, A
C 21	15.2	69.1	25	7	US-11-136-527-311448 Sequence 311448, A
C 22	15.2	69.1	448	7	US-11-123-896-406 Sequence 406, App
C 23	15.2	69.1	600	7	US-11-136-527-6599 Sequence 6599, Ap

C 24	15.2	69.1	1870	6	US-10-750-185-38563 Sequence 38563, A
C 25	15.2	69.1	1870	6	US-10-750-623-38563 Sequence 38563, A
C 26	15.2	69.1	3328	7	US-11-136-527-2503 Sequence 2503, Ap
C 27	15	68.2	1400	7	US-11-136-527-4366 Sequence 4366, Ap
C 28	15	68.2	2746	7	US-11-136-527-270 Sequence 270, App
C 29	14.8	67.3	25	7	US-11-136-527-311434 Sequence 311434, A
C 30	14.8	67.3	1509	6	US-10-454-437-211 Sequence 211, App
C 31	14.8	67.3	4091	6	US-10-750-185-32344 Sequence 32344, A
C 32	14.8	67.3	4091	6	US-10-750-623-32344 Sequence 32344, A
C 33	14.8	67.3	182303	7	US-11-121-086-45 Sequence 45, Appl
C 34	14.6	66.4	408	6	US-10-793-626-2675 Sequence 2675, Ap
C 35	14.6	66.4	600	6	US-10-750-185-3617 Sequence 3617, Ap
C 36	14.6	66.4	600	6	US-10-750-623-3617 Sequence 3617, Ap
C 37	14.6	66.4	1161	6	US-10-793-626-2299 Sequence 2299, Ap
C 38	14.6	66.4	1398	7	US-11-133-360-2 Sequence 2, Appl
C 39	14.6	66.4	1401	7	US-11-133-360-6 Sequence 6, Appl
C 40	14.6	66.4	1401	7	US-11-133-360-8 Sequence 8, Appl
C 41	14.6	66.4	1951	6	US-10-750-185-64757 Sequence 64757, A
C 42	14.6	66.4	1951	6	US-10-750-623-64757 Sequence 64757, A
C 43	14.6	66.4	2083	7	US-11-167-856-15 Sequence 15, Appl
C 44	14.6	66.4	2290	6	US-10-750-185-44532 Sequence 44532, A
C 45	14.6	66.4	2290	6	US-10-750-623-44532 Sequence 44532, A

ALIGNMENTS

RESULT 1
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; Sequence 53745, Application US/10750185
; Publication No. US200502603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10750,185
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 53745
; LENGTH: 1107
; TYPE: DNA
; ORGANISM: Bovine 19866881030294
US-10-750-185-53745
Query Match 76.4%; Score 16.8; DB 6; Length 1107;
Best Local Similarity 90.0%; Pred. No. 45;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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DB 800 TAACCACCATCGATCAATG 781
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; Sequence 53745, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen

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; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10/750,623
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 53745
; LENGTH: 1107
; TYPE: DNA
; ORGANISM: Bovine 19866881030294
US-10-750-623-53745

Query Match      76.4%; Score 16.8; DB 6; Length 1107;
Best Local Similarity 90.0%; Pred. No. 45;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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RESULT 3
US-11-133-360-4
; Sequence 4, Application US/11133360
; Publication No. US20050287172A1
; GENERAL INFORMATION:
; APPLICANT: Medimmune Vaccines, Inc et al.
; TITLE OF INVENTION: INFLUENZA HEMAGGLUTININ AND NEURAMINIDASE VARIANTS
; FILE REFERENCE: FL260
; CURRENT APPLICATION NUMBER: 2005-05-20
; CURRENT FILING DATE: 2005-05-20
; PRIOR APPLICATION NUMBER: 60/574,553
; PRIOR FILING DATE: 2004-05-25
; PRIOR APPLICATION NUMBER: 60/657,554
; PRIOR FILING DATE: 2005-02-28
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 4
; LENGTH: 1458
; TYPE: DNA
; ORGANISM: Influenza virus
US-11-133-360-4

Query Match      73.6%; Score 16.2; DB 7; Length 1458;
Best Local Similarity 85.7%; Pred. No. 90;
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RESULT 4
US-10-750-185-46643
; Sequence 46643, Application US/10750185
; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10/750,185
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
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; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 46643
; LENGTH: 1786
; TYPE: DNA
; ORGANISM: Bovine 19866881022693
US-10-750-185-46643

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Best Local Similarity 85.7%; Pred. No. 92;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

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Db 882 TGACCACCATCGCATCACTGT 902

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; Sequence 46643, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10/750,623
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 46643
; LENGTH: 1786
; TYPE: DNA
; ORGANISM: Bovine 19866881022693
US-10-750-623-46643

Query Match      73.6%; Score 16.2; DB 6; Length 1786;
Best Local Similarity 85.7%; Pred. No. 92;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGT 21
Db 882 TGACCACCATCGCATCACTGT 902

RESULT 6
US-11-128-061-3932
; Sequence 3932, Application US/11128061
; Publication No. US20060003958A1
; GENERAL INFORMATION:
; APPLICANT: Melville, Mark W.
; APPLICANT: Charlebois, Timothy S.
; APPLICANT: Mounts, William M.
; APPLICANT: Hann, Louane E.
; APPLICANT: Sinacore, Martin S.
; APPLICANT: Leonard, Mark W.
; APPLICANT: Brown, Eugene L.
; APPLICANT: MILLER, Christopher P.
; TITLE OF INVENTION: NOVEL POLYNUCLEOTIDES RELATED TO OLIGONUCLEOTIDE ARRAYS
; FILE REFERENCE: 01997.027701
; CURRENT APPLICATION NUMBER: US/11/128,061
; CURRENT FILING DATE: 2005-05-11
; PRIOR APPLICATION NUMBER: US 60/570,425
; PRIOR FILING DATE: 2004-05-11
; NUMBER OF SEQ ID NOS: 7285
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Title: US-10-754-446-2

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8: /cgn2_6/ptodata/1/ina/RE COMB.seq.*
9: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	22	100.0	2051	3	US-09-949-016-1712 Sequence 1712, Ap
C 2	22	100.0	15353	3	US-09-949-016-13454 Sequence 13454, A
C 3	16	72.7	1650	3	US-09-107-532A-3186 Sequence 3186, Ap
C 4	15.8	71.8	537	3	US-09-489-039A-5168 Sequence 5168, Ap
C 5	15.8	71.8	572	3	US-09-533-559-6112 Sequence 6112, Ap
C 6	15.8	71.8	601	3	US-09-949-016-79854 Sequence 79854, A
C 7	15.8	71.8	601	3	US-09-949-016-83923 Sequence 83923, A
C 8	15.8	71.8	1614	2	US-08-867-941-10 Sequence 10, Appl
C 9	15.8	71.8	1614	2	US-08-867-941-10 Sequence 10, Appl
C 10	15.8	71.8	1623	2	US-08-867-941-5 Sequence 5, Appl
C 11	15.8	71.8	1623	2	US-08-867-941-5 Sequence 5, Appl
C 12	15.8	71.8	1635	3	US-09-540-236-840 Sequence 840, App
C 13	15.8	71.8	4657	3	US-09-254-325-1 Sequence 1, Appl
C 14	15.8	71.8	5643	3	US-09-079-415-5 Sequence 5, Appl
C 15	15.8	71.8	5643	3	US-08-750-458A-1 Sequence 1, Appl
C 16	15.8	71.8	7641	2	US-08-867-941-6 Sequence 6, Appl
C 17	15.8	71.8	7641	3	US-09-074-658-6 Sequence 6, Appl
C 18	15.8	71.8	7650	2	US-08-867-941-1 Sequence 1, Appl
C 19	15.8	71.8	7650	3	US-09-074-658-1 Sequence 1, Appl
C 20	15.8	71.8	4322	3	US-10-024-396-13 Sequence 13, Appl
C 21	15.8	71.8	49617	3	US-09-596-002-28 Sequence 28, Appl
C 22	15.8	71.8	90472	3	US-09-949-016-14038 Sequence 14038, A
C 23	15.8	71.8	96340	3	US-09-949-016-15863 Sequence 15863, A
C 24	15.8	71.8	455726	3	US-09-949-016-14157 Sequence 14157, A

C 25	15.8	71.8	481115	3	US-09-949-016-11940 Sequence 11940, A
C 26	15.6	70.9	762	3	US-09-252-991A-3481 Sequence 3481, Ap
C 27	15.6	70.9	816	3	US-09-489-039A-4176 Sequence 4176, Ap
C 28	15.6	70.9	834	3	US-09-489-039A-2174 Sequence 2174, Ap
C 29	15.6	70.9	1203	3	US-09-252-991A-3468 Sequence 3468, Ap
C 30	15.6	70.9	1269	3	US-09-252-991A-3470 Sequence 3470, Ap
C 31	15.6	70.9	1428	3	US-09-248-796A-3428 Sequence 3428, Ap
C 32	15.6	70.9	1697	3	US-09-270-767-10255 Sequence 10255, A
C 33	15.6	70.9	1876	3	US-08-192-271-8 Sequence 8, Appl
C 34	15.6	70.9	1876	6	PCT-US94-01572-8 Sequence 8, Appl
C 35	15.6	70.9	2231	3	US-09-949-016-4932 Sequence 4932, Ap
C 36	15.6	70.9	2231	3	US-09-949-016-4933 Sequence 4933, Ap
C 37	15.6	70.9	3041	3	US-10-126-120-1 Sequence 1, Appl
C 38	15.6	70.9	4081	3	US-08-939-774A-1 Sequence 1, Appl
C 39	15.6	70.9	4558	2	US-08-309-512-2 Sequence 2, Appl
C 40	15.6	70.9	4558	6	PCT-US92-08756A-2 Sequence 2, Appl
C 41	15.6	70.9	7400	2	US-07-674-852-1 Sequence 1, Appl
C 42	15.6	70.9	7400	3	US-08-473-185-1 Sequence 1, Appl
C 43	15.6	70.9	7400	3	US-09-171-387-3 Sequence 3, Appl
C 44	15.6	70.9	113379	3	US-09-949-016-17561 Sequence 17561, A
C 45	15.6	70.9	113379	3	US-09-949-016-17562 Sequence 17562, A

ALIGNMENTS

RESULT 1

US-09-949-016-1712/c
; Sequence 1712, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1712
; LENGTH: 2051
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-1712

Query Match 100.0%; Score 22; DB 3; Length 2051;

Best Local Similarity 100.0%; Pred. No. 0.24; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22

DB 693 TAACCACCATCGGATCAATGTC 672

RESULT 2

US-09-949-016-13454/c
; Sequence 13454, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20

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; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13454
; LENGTH: 15353
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13454

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Best Local Similarity 100.0%; Pred. No. 0.3;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAATGTC 22
Db 6298 TAACCACCATCGGATCAATGTC 6277

RESULT 3
US-09-107-532A-3186/c
; Sequence 3186, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Walcham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 3186:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1650 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...1650
; SEQUENCE DESCRIPTION: SEQ ID NO: 3186:
US-09-107-532A-3186
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QY 2 AACCCACCATCGGATCA 17
Db 578 AACCCACCATCGGATCA 563

RESULT 4
US-09-489-039A-5168/c
; Sequence 5168, Application US/09489039A
; Patent No. 6610836
; GENERAL INFORMATION:
; APPLICANT: Gary Breton et. al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO KLEBSIELLA
; PNEUMONIAE FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2004001
; CURRENT APPLICATION NUMBER: US/09/489,039A
; CURRENT FILING DATE: 2000-01-27
; PRIOR APPLICATION NUMBER: US 60/117,747
; PRIOR FILING DATE: 1999-01-29
; NUMBER OF SEQ ID NOS: 14342
; SEQ ID NO 5168
; LENGTH: 537
; TYPE: DNA
; ORGANISM: Klebsiella pneumoniae
US-09-489-039A-5168

Query Match      71.8%; Score 15.8; DB 3; Length 537;
Best Local Similarity 89.5%; Pred. No. 2.4e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TAACCACCATCGGATCAAT 19
Db 151 TAACCACCATCGATTCAT 133
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RESULT 5
US-09-533-559-6112
; Sequence 6112, Application US/09533559
; Patent No. 6902887
; GENERAL INFORMATION:
; APPLICANT: Randy M. Berka
; APPLICANT: Michael W. Rey
; APPLICANT: Jeffrey R. Shuster
; APPLICANT: Sakari Kauppinen
; APPLICANT: Ib Groth Clausen
; APPLICANT: Peter Bjarke Olsen
; TITLE OF INVENTION: Methods For Monitoring Multiple Gene
; EXPRESSION
; FILE REFERENCE: 5849.200-US
; CURRENT APPLICATION NUMBER: US/09/533,559
; CURRENT FILING DATE: 2000-03-22
; EARLIER APPLICATION NUMBER: 09/273,623
; EARLIER FILING DATE: 1999-03-22
; NUMBER OF SEQ ID NOS: 7860
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 6112
; LENGTH: 572
; TYPE: DNA
; ORGANISM: Aspergillus oryzae
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(572)
; OTHER INFORMATION: n = A,T,C or G
US-09-533-559-6112
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Best Local Similarity 89.5%; Pred. No. 2.5e+02;
Matches 17; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:49:05 ; Search time 736.68 Seconds
(without alignments)
1157.425 Million cell updates/sec

Title: US-10-754-446-1

Perfect score: 15

Sequence: 1 agcggcgagactca 15

Scoring table:
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Gapop 10.0 , Gapext 1.0

Searched: 5883141 seqs, 28421725653 residues

Total number of hits satisfying chosen parameters: 11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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2: gb.in.*
3: gb.env.*
4: gb.om.*
5: gb.ov.*
6: gb.pat.*
7: gb.ph.*
8: gb.pr.*
9: gb.ro.*
10: gb.sts.*
11: gb.sv.*
12: gb.un.*
13: gb.vi.*
14: gb.htg.*
15: gb.pl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	15	100.0	21081	2	AF106590 Caenorhab
4	15	100.0	36859	2	AF003141 Caenorhab
5	15	100.0	103303	14	AY339213 Takifugu
6	15	100.0	110000	1	Continuation (4 of
7	15	100.0	155645	14	AC021153 Homo sapi
8	15	100.0	166981	14	AC157751 Callithri
9	15	100.0	173126	8	AC008878 Homo sapi
10	15	100.0	180213	14	AC150812 Callithri
11	15	100.0	197202	14	AC153077 Callithri
12	15	100.0	208232	9	AC149085 Mus muscu
13	15	100.0	221553	9	AC149282 Mus muscu
14	15	100.0	228698	14	AC127479 Mus muscu
15	15	100.0	237653	14	AC073760 Mus muscu
16	15	100.0	291000	1	SC0939105 Streptomy
17	14	93.3	563	15	AJ836913 Arabidops
18	14	93.3	789	9	AF296282 Mus muscu

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c 20	14	93.3	1250	9	S70756S1	S70756 Tlx-1-homoe
c 21	14	93.3	1950	6	AR622581	Sequence
c 22	14	93.3	2262	9	AB127583	Mus muscu
c 23	14	93.3	2428	8	AK096823	Homo sapi
c 24	14	93.3	2685	9	BC059941	Mus muscu
c 25	14	93.3	3238	5	AF427524	Danio rer
c 26	14	93.3	4102	6	CQ986467	Sequence
c 27	14	93.3	10246	9	MMU251787	AJ251787 Mus muscu
c 28	14	93.3	10586	1	AE004991	Halobacte
c 29	14	93.3	10696	1	AE012476	Xanthomon
c 30	14	93.3	20074	8	HSA9794	Homo sapi
c 31	14	93.3	20232	6	AR619802	Sequence
c 32	14	93.3	45653	5	EX323016	BX323016 Zebrafish
c 33	14	93.3	51099	8	AL357395	AJ357395 Human DNA
c 34	14	93.3	53425	14	AY211534	Zea mays
c 35	14	93.3	89015	14	AC008458	Homo sapi
c 36	14	93.3	94092	14	AC155398	Zea mays
c 37	14	93.3	110000	1	AE016822_15	Continuation (16 o
c 38	14	93.3	110000	1	AE017282_18	Continuation (19 o
c 39	14	93.3	110000	1	AE017282_29	Continuation (30 o
c 40	14	93.3	110000	1	AE017340_16	Continuation (17 o
c 41	14	93.3	110000	1	AP006618_44	Continuation (45 o
c 42	14	93.3	110000	1	AP006618_45	Continuation (46 o
c 43	14	93.3	110000	1	AP006618_58	Continuation (59 o
c 44	14	93.3	110000	1	AY596297_08	Continuation (9 of
c 45	14	93.3	110000	1	AY596297_28	Continuation (29 o

ALIGNMENTS

RESULT 1
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LOCUS Homo sapiens mucopolin 1 (MCOLN1) gene, exons 3 and 4.
ACCESSION AF305574
VERSION AF305574.1 GI:11991200
KEYWORDS 3 of 8
SEGMENT
SOURCE Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
Hominidae; Homo.
REFERENCE 1 (bases 1 to 790)
AUTHORS Bargal, R., Avidan, N., Ben-Asher, E., Olender, Z., Zeigler, M.,
Franklin, A., Raas-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
TITLE Identification of the gene causing mucopolidiosis type IV
JOURNAL Nat. Genet. 26 (1), 118-123 (2000)
PUBMED 10973263
REFERENCE 2 (bases 1 to 790)
AUTHORS Bargal, R., Avidan, N., Ben-Asher, E., Olender, A., Zeigler, M.,
Franklin, A., Raas-Rothschild, A., Glusman, G., Lancet, D. and Bach, G.
TITLE Direct Submission
JOURNAL Submitted (13-SEP-2000) Molecular Genetics, The Weizmann Institute
of Science, P. O. Box 26, Rehovot 76100, Israel
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589..757
/gene="MCOLN1"
/number=4
exon
exon
ORIGIN
Query Match 100.0% Score 15; DB 8; Length 790;
Best Local Similarity 100.0%; Pred. No. 7.3e+03; Indels 0; Gaps 0;
Matches 15; Conservative 0; Mismatches 0;

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: January 15, 2006, 15:55:50 ; Search time 1677.48 Seconds
(without alignments)
418.369 Million cell updates/sec

Title: US-10-754-446-1

Perfect score: 15

Sequence: 1 agcgggcccggactca 15

Scoring table: IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters: 82156650

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST.*

1: gb_est1.*

2: gb_est2.*

3: gb_est3.*

4: gb_hc.*

5: gb_est4.*

6: gb_est5.*

7: gb_est6.*

8: gb_est7.*

9: gb_gse1.*

10: gb_gse2.*

11: gb_gse3.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	15	100.0	301	5	BY345580 BY345580
C 2	15	100.0	331	5	BY345176 BY345176
C 3	15	100.0	338	5	BY102360 BY102360
C 4	15	100.0	375	5	BY087147 BY087147
C 5	15	100.0	375	5	C08554 C08554
C 6	15	100.0	378	5	BY299457 BY299457
C 7	15	100.0	379	5	C71190 C71190
C 8	15	100.0	387	9	CC423137 PUEH772TD
C 9	15	100.0	469	1	AV631303 AV631303
C 10	15	100.0	480	5	BY249487 BY249487
C 11	15	100.0	502	7	CK583489 CK583489
C 12	15	100.0	519	7	CK583423 CK583423
C 13	15	100.0	568	7	CK577649 CK577649
C 14	15	100.0	624	3	BY726669 BY726669
C 15	15	100.0	638	1	AU200865 AU200865
C 16	15	100.0	779	10	CG116594 CG116594
C 17	15	100.0	786	10	AG602966 AG602966
C 18	15	100.0	797	7	CN200405 CN200405
C 19	15	100.0	832	8	CK892033 CK892033
C 20	15	100.0	962	6	CA464646 CA464646
C 21	15	100.0	1040	3	BQ278046 BQ278046
C 22	14	93.3	194	3	BQ082114 BQ082114

23	14	93.3	208	2	BB568361
24	14	93.3	287	9	AZ121792
c 25	14	93.3	301	1	A1029662
26	14	93.3	306	2	BB419855
27	14	93.3	350	8	W66660
c 28	14	93.3	354	5	BY312288
29	14	93.3	356	8	W99177
30	14	93.3	360	5	C45921
31	14	93.3	374	5	C43796
c 32	14	93.3	389	8	DN368435
33	14	93.3	402	11	DR18114S
c 34	14	93.3	409	5	BY237451
35	14	93.3	436	9	AZ251529
36	14	93.3	455	1	AA050210
37	14	93.3	462	5	BX527893
c 38	14	93.3	469	7	CJ034986
39	14	93.3	471	8	W99993
c 40	14	93.3	473	8	DN367700
c 41	14	93.3	484	5	BX635951
c 42	14	93.3	487	10	CG732852
43	14	93.3	489	8	DN481676
44	14	93.3	489	10	CG465566
c 45	14	93.3	491	2	BE943858

ALIGNMENTS

RESULT 1
BY345580/c 301 bp mRNA linear EST 12-DEC-2002
LOCUS
DEFINITION BY345580 RIKEN full-length enriched, whole joints Mus musculus cDNA
clone L730015A09 5', mRNA sequence.
ACCESSION BY345580.1 GI:26575068
VERSION
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muroidae; Muridae; Murinae; Mus.
1 (bases 1 to 301)

Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Osato, N., Salto, R., Suzuki, H., Yamada, H.,
Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A.,
Schonbach, C., Gojohori, T., Baldarelli, R., Hill, D. P., Bult, C.,
Hume, D. A., Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H.,
Batalov, S., Beisel, K. W., Blake, J. A., Bradt, D., Brusic, V.,
Chothia, C., Corbani, L. E., Cousins, S., Dalla, E., Dragani, T. A.,
Fletcher, C. F., Forrest, A., Frazer, K. S., Gaasterland, T.,
Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S.,
Gustincich, S., Hirokawa, N., Jackson, I. J., Jarvis, E. D., Kanai, A.,
Kawaji, H., Kawasawa, Y., Kedzierski, R. M., King, B. L., Konagaya, A.,
Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R.,
Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T.,
Numata, K., Okido, T., Pavan, W. J., Pertea, G., Pesole, G.,
Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramachandran, S.,
Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring, B. Z., Ringwald, M.,
Sandelin, A., Schneider, C., Semple, C. A., Setou, M., Shimada, K.,
Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale, R. D., Tonita, M.,
Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y.,
Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagiya, M., Yang, I.,
Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P.,
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Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K.,
Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y.,
Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K.,
Shingawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Landet, E. S.,
Rogers, J., Birney, E. and Hayashizaki, Y.
Analysis of the mouse transcriptome based on functional annotation
of 60,770 full-length cDNAs
NATURE 420, 563-573 (2002)
JOURNAL
PUBMED
12466851

COMMENT

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Email: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>
Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S.,
Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H.,
Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R.,
Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K.,
Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and
Hayashizaki, Y. Direct Submission
Computational Analysis of Full-Length Mouse cDNAs Compared with
Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
cDNA library was prepared and sequenced in Mouse Genome
Encyclopedia Project of Genome Exploration Research Group in Riken
Genomic Sciences Center and Genome Science Laboratory in RIKEN.
Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Tissues were provided by Vassilis Aidinis (Biomedical Sciences
Research Center 'Al. Fleming' Institute of Immunology 14-16 Al.
Fleming street 16672 Vari,Greece) whose assistance we gratefully
acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.

FEATURES

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LOCUS

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ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

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Mus musculus
Mus musculus (house mouse)
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Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
Sciurognathi; Muridae; Muridae; Murinae; Mus.

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REFERENCE

AUTHORS

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1 (bases 1 to 331)
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamanaka, I.,
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Kawai, H., Kawasawa, Y., Kedzierski, R.M., King, B.L., Konagaya, A.,
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Contact: Yoshihide Hayashizaki
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The Institute of Physical and Chemical Research (RIKEN)
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prepare full-length cDNA libraries for rapid discovery of new
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Division of Experimental Animal Research in Riken contributed to
prepare mouse tissues.
Tissues were provided by Vassilis Aidinis (Biomedical Sciences
Research Center 'Al. Fleming' Institute of Immunology 14-16 Al.
Fleming street 16672 Vari,Greece) whose assistance we gratefully
acknowledge.
Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.

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OM nucleic - nucleic search, using sw model

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(without alignments)
491.207 Million cell updates/sec

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14: Geneseq2005s.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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c 25	13.4	89.3	616	13 ADU52117	Adu52117 Fusarium
c 26	13.4	89.3	616	14 ADZ90120	Adz90120 Fusarium
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30	13.4	89.3	681	9 ACD10983	Acd10983 Human col
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33	13.4	89.3	940	4 AAS01548	Aas01548 Human sec
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43	13.4	89.3	1676	14 ADY17906	Ady17906 DNA encod
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ALIGNMENTS

RESULT 1

AEB28957
ID AEB28957 standard; DNA; 15 BP.

XX AC AEB28957;

XX DT 22-SEP-2005 (first entry)

XX DE Human MCOLN1 gene PCR primer IVS-F SEQ ID NO:1.

XX KW mucolipidosis IV; high throughput screening; mucolipin-1; MCOLN1; PCR; primer; ss.

XX OS Homo sapiens.

XX PN US2005153300-A1.

XX PD 14-JUL-2005.

XX PF 09-JAN-2004; 2004US-00754446.

XX PR 09-JAN-2004; 2004US-00754446.

XX (QUES-) QUEST DIAGNOSTICS INC.

XX PI Sun W, Hantash F;

XX DR WPI; 2005-521160/53.

XX PT Diagnosing mucolipidosis (ML) IV mutant sequence in nucleic acids, by amplifying the nucleic acid, detecting amplified product with labeled oligonucleotide probes via a change in fluorescence which indicates the presence of an ML IV mutant.

XX PS Claim 1; SEQ ID NO 1; 15pp; English.

XX CC The invention relates to a method (M1) for detecting the presence of mucolipidosis IV (ML IV) mutant sequences in a nucleic acid. (M1) involves contacting the nucleic acid with oligonucleotide primers and probes, conducting amplification by temperature cycling and monitoring the accumulation of amplified nucleic acid by detecting an increase in donor fluorophore fluorescence or decrease in acceptor fluorophore

CC fluorescence, which indicates the presence of ML IV mutant sequence. Also
CC described: (1) an oligonucleotide (ON1) comprising a sequence
CC complementary to the coding or non-coding strand of the mucopolin-1
CC (MCOLN1) gene; and (2) a kit (K1) for amplifying sequences of ML IV
CC sequences, comprising ON1. (M1) is useful for determining the presence of
CC ML IV mutant sequence in a nucleic acid, and/or wild-type sequences. (M1)
CC is useful for genetic counseling of individuals at risk for ML IV or at
CC risk to having offspring afflicted with ML IV. The MCOLN1 gene is located
CC on chromosome 19p13. The present sequence represents a PCR primer for the
CC MCOLN1 gene, which is used in the exemplification of the present
CC invention.

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ID AAK73827 standard; DNA; 11365 BP.

AC AAK73827;

DT 07-NOV-2001 (first entry)

XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:28639.

XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;
XX Cytostatic; Gene therapy; vaccine; metastasis; db.

OS Homo sapiens.

XX WO200157182-A2.

XX 09-AUG-2001.

XX 17-JAN-2001; 2001WO-US0001354.

XX 31-JAN-2000; 2000US-0179065P.

PR 04-FEB-2000; 2000US-0180628P.

PR 24-FEB-2000; 2000US-0184664P.

PR 02-MAR-2000; 2000US-0186350P.

PR 16-MAR-2000; 2000US-0189874P.

PR 17-MAR-2000; 2000US-0190076P.

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PR 19-MAY-2000; 2000US-0205515P.

PR 07-JUN-2000; 2000US-0209467P.

PR 28-JUN-2000; 2000US-0214866P.

PR 30-JUN-2000; 2000US-0215135P.

PR 07-JUL-2000; 2000US-0216647P.

PR 07-JUL-2000; 2000US-0216880P.

PR 11-JUL-2000; 2000US-0217487P.

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PR 14-AUG-2000; 2000US-0224518P.

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PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.

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OM nucleic - nucleic search, using sw model

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(without alignments)
342.389 Million cell updates/sec

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Perfect score: 15

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Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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5	14	93.3	432	7	US-10-430-201-1048
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7	14	93.3	527	5	US-10-027-632-130943
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10	14	93.3	527	6	US-10-027-632-130944
11	14	93.3	606	7	US-10-767-701-524
12	14	93.3	837	6	US-10-369-493-37513
13	14	93.3	858	8	US-10-425-115-162360
14	14	93.3	4102	8	US-10-602-494-17
15	14	93.3	19265	8	US-10-719-993-6839
16	13.4	89.3	63	7	US-10-362-906-3
17	13.4	89.3	252	6	US-10-029-386-26242
18	13.4	89.3	303	3	US-09-974-300-5829
19	13.4	89.3	329	8	US-10-425-115-122740
20	13.4	89.3	399	9	US-10-779-543-12575
21	13.4	89.3	438	7	US-10-437-963-5136
22	13.4	89.3	443	8	US-10-425-115-29829
23	13.4	89.3	483	4	US-09-925-065A-755757

C 24	13.4	89.3	499	4	US-09-925-065A-762110	Sequence 762110, A
25	13.4	89.3	556	6	US-10-029-386-12542	Sequence 12542, A
C 26	13.4	89.3	610	5	US-10-027-632-99325	Sequence 99325, A
C 27	13.4	89.3	610	8	US-10-027-632-99325	Sequence 99325, A
C 28	13.4	89.3	616	8	US-10-653-047-599	Sequence 599, App
29	13.4	89.3	638	8	US-10-425-115-140595	Sequence 140595, A
30	13.4	89.3	645	5	US-10-027-632-28813	Sequence 28813, A
31	13.4	89.3	645	5	US-10-027-632-28814	Sequence 28814, A
32	13.4	89.3	645	6	US-10-027-632-28813	Sequence 28813, A
33	13.4	89.3	645	6	US-10-027-632-28814	Sequence 28814, A
34	13.4	89.3	648	6	US-10-369-493-39498	Sequence 39498, A
35	13.4	89.3	648	6	US-10-369-493-39877	Sequence 39877, A
36	13.4	89.3	651	6	US-10-369-493-39129	Sequence 39129, A
37	13.4	89.3	681	5	US-10-001-883-38	Sequence 38, Appl
38	13.4	89.3	861	6	US-10-156-761-2634	Sequence 2634, Ap
39	13.4	89.3	968	8	US-10-653-047-7533	Sequence 7533, Ap
40	13.4	89.3	984	6	US-10-156-761-1305	Sequence 1305, Ap
C 41	13.4	89.3	1052	9	US-10-450-763-20158	Sequence 20158, A
C 42	13.4	89.3	1152	6	US-10-369-493-46700	Sequence 46700, A
43	13.4	89.3	1353	6	US-10-156-761-4728	Sequence 4728, Ap
44	13.4	89.3	1450	8	US-10-425-115-140634	Sequence 140634, A
45	13.4	89.3	1500	10	US-11-097-143-35051	Sequence 35051, A

ALIGNMENTS

RESULT 1
US-10-754-446-1
; Sequence 1, Application US/10754446
; Publication No. US20050153300A1
; GENERAL INFORMATION:
; APPLICANT: SUN, WEIMIN
; APPLICANT: HANTASH, FERAS
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR THE DETECTION OF
; TITLE OF INVENTION: MUCOLIPIDOSIS IV MUTATIONS
; FILE REFERENCE: 034827-2301
; CURRENT APPLICATION NUMBER: US/10/754,446
; CURRENT FILING DATE: 2004-01-09
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: Patent In Ver. 3.2
; SEQ ID NO 1
; LENGTH: 15
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Primer
US-10-754-446-1

Query Match 100.0%; Score 15; DB 9; Length 15;
Best Local Similarity 100.0%; Pred. No. 4e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 AGCGGCGCGACTCA 15
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Db 1 AGCGGCGCGACTCA 15

RESULT 2
US-10-425-115-103385
; Sequence 103385, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated with
; TITLE OF INVENTION: Plants
; FILE REFERENCE: 38-21(53222)B
; CURRENT APPLICATION NUMBER: US/10/425,115
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326

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; SEQ ID NO 103385
; LENGTH: 356
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)..(356)
; OTHER INFORMATION: unsure at all n locations
; FEATURE:
; OTHER INFORMATION: Clone ID: MRT4577_25792C.1
US-10-425-115-103385

Query Match      100.0%; Score 15; DB 8; Length 356;
Best Local Similarity 100.0%; Pred. No. 1.8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 AGCGGGCGGACTCA 15
Db      255 AGCGGGCGGACTCA 269
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RESULT 3
US-09-851-494B-1
; Sequence 1, Application US/09851494B
; Publication No. US20030064363A1
; GENERAL INFORMATION:
; APPLICANT: ML4 Foundation
; APPLICANT: Goldin, Ehud
; APPLICANT: Staugenhaupt, Susan A.
; APPLICANT: Sun, Mei
; APPLICANT: Agierno, James S.
; TITLE OF INVENTION: A Gene Encoding A New TRP Channel is Mutated in Mucopolipidosis IV
; FILE REFERENCE: 3394/1H557U51
; CURRENT APPLICATION NUMBER: US/09/851.494B
; CURRENT FILING DATE: 2002-07-12
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 13270
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-851-494B-1

Query Match      100.0%; Score 15; DB 3; Length 13270;
Best Local Similarity 100.0%; Pred. No. 71;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      5495 AGCGGGCGGACTCA 5509
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RESULT 4
US-10-719-993-29684/c
; Sequence 29684, Application US/10719993
; Publication No. US20040265849A1
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele et al.
; TITLE OF INVENTION: GENETIC POLYMORPHISMS ASSOCIATED WITH
; FILE REFERENCE: ALZHEIMER'S DISEASE, METHODS OF DETECTION AND USES THEREOF
; CURRENT APPLICATION NUMBER: US/10/719,993
; CURRENT FILING DATE: 2003-11-24
; NUMBER OF SEQ ID NOS: 55342
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 29684
; LENGTH: 201
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-719-993-29684

Query Match      93.3%; Score 14; DB 8; Length 201;
Best Local Similarity 100.0%; Pred. No. 7.1e+02;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db      159 GCGGGCGGACTCA 146
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RESULT 5
US-10-430-201-1048
; Sequence 1048, Application US/10430201
; Publication No. US20040162679A1
; GENERAL INFORMATION:
; APPLICANT: Li, Linheng
; TITLE OF INVENTION: Method for Predicting Gene Potential and Cell Commitment
; FILE REFERENCE: 40716 (1p-010)
; CURRENT APPLICATION NUMBER: US/10/430,201
; CURRENT FILING DATE: 2003-05-05
; PRIOR APPLICATION NUMBER: US 60/370,114
; PRIOR FILING DATE: 2002-05-03
; NUMBER OF SEQ ID NOS: 4879
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 1048
; LENGTH: 432
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (212)..(212)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
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; OTHER INFORMATION: n is a, c, g, or t
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GenCore version 5.1.6
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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	13.4	89.3	182314	7	US-11-112-908-45
4	13	86.7	1321	6	US-10-750-185-49780
5	13	86.7	1321	6	US-10-750-623-49780
6	13	86.7	15876	7	US-11-052-554A-660
C 7	12.4	82.7	65	6	US-10-310-914A-7133
C 8	12.4	82.7	65	6	US-10-310-914A-8862
9	12.4	82.7	126	6	US-10-467-657-867
10	12.4	82.7	153	6	US-10-467-657-735
11	12.4	82.7	162	6	US-10-467-657-743
C 12	12.4	82.7	201	6	US-10-995-561-11825
C 13	12.4	82.7	201	6	US-10-995-561-61260
C 14	12.4	82.7	201	6	US-10-995-561-82266
C 15	12.4	82.7	558	7	US-11-212-443-49
C 16	12.4	82.7	600	6	US-10-750-185-21924
C 17	12.4	82.7	600	6	US-10-750-623-21924
C 18	12.4	82.7	600	7	US-11-136-527-6844
C 19	12.4	82.7	639	7	US-11-212-443-47
C 20	12.4	82.7	660	7	US-11-212-443-45
C 21	12.4	82.7	696	6	US-10-467-657-3351
C 22	12.4	82.7	960	6	US-10-750-185-63567
23	12.4	82.7	960	6	US-10-750-623-63567

C 24	12.4	82.7	999	6	US-10-750-185-53582	Sequence 53582, A
C 25	12.4	82.7	999	6	US-10-750-623-53582	Sequence 53582, A
C 26	12.4	82.7	1369	7	US-11-128-061-3328	Sequence 3328, Ap
C 27	12.4	82.7	1400	7	US-11-136-527-1821	Sequence 1821, Ap
C 28	12.4	82.7	1465	6	US-10-750-185-62827	Sequence 62827, A
C 29	12.4	82.7	1465	6	US-10-750-623-62827	Sequence 62827, A
C 30	12.4	82.7	1581	6	US-10-750-185-64370	Sequence 64370, A
C 31	12.4	82.7	1581	6	US-10-750-623-64370	Sequence 64370, A
C 32	12.4	82.7	1604	7	US-11-192-967-7	Sequence 7, Appli
C 33	12.4	82.7	1604	7	US-11-193-715-7	Sequence 7, Appli
C 34	12.4	82.7	1713	7	US-11-136-527-2748	Sequence 2748, Ap
C 35	12.4	82.7	1853	6	US-10-967-527A-1	Sequence 1, Appli
36	12.4	82.7	1938	6	US-10-967-527A-29	Sequence 29, Appl
37	12.4	82.7	1979	6	US-10-967-527A-31	Sequence 31, Appl
38	12.4	82.7	2232	6	US-10-986-501-79	Sequence 79, Appl
39	12.4	82.7	2879	7	US-11-124-368A-166	Sequence 166, App
40	12.4	82.7	3635	7	US-11-136-527-2101	Sequence 2101, Ap
41	12.4	82.7	4734	7	US-11-136-527-2518	Sequence 2518, Ap
C 42	12.4	82.7	5498	7	US-11-192-967-1	Sequence 1, Appli
C 43	12.4	82.7	5498	7	US-11-193-715-1	Sequence 1, Appli
C 44	12.4	82.7	6719	6	US-10-995-561-456	Sequence 456, App
45	12.4	82.7	9903	7	US-11-052-554A-517	Sequence 517, App

ALIGNMENTS

RESULT 1

US-10-310-914A-1387400
; Sequence 1387400, Application US/10310914A
; Publication No. US20060003322A1
; GENERAL INFORMATION:
; APPLICANT: Bentwich, Isaac
; APPLICANT: Shiller, Kvuzat
; TITLE OF INVENTION: Bioinformatically detectable group of novel regulatory genes and
; FILE REFERENCE: 06087.0200.CPUS01
; CURRENT APPLICATION NUMBER: US/10/310,914A
; CURRENT FILING DATE: 2002-12-06
; NUMBER OF SEQ ID NOS: 1388402
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 1387400
; LENGTH: 20
; TYPE: RNA
; ORGANISM: Human
US-10-310-914A-1387400

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Best Local Similarity 86.7%; Pred. No. 8.6e+02;
Matches 13; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 AGCGGGCGGACTCA 15

Db 2 AGCGGGCGGACTCA 16

RESULT 2

US-11-121-849-654079
; Sequence 654079, Application US/11121849
; Publication No. US20050272080A1
; GENERAL INFORMATION:
; APPLICANT: John Palma
; TITLE OF INVENTION: Methods of Genetic Analysis of Formalin Fixed Paraffin Embedded S
; FILE REFERENCE: 3684.1
; CURRENT APPLICATION NUMBER: US/11/121,849
; CURRENT FILING DATE: 2005-05-03
; PRIOR FILING DATE: 2004-05-03
; NUMBER OF SEQ ID NOS: 673904
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 654079
; LENGTH: 25

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; TYPE: DNA
; ORGANISM: Homo sapien
US-11-121-849-654079

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Best Local Similarity 93.3%; Pred. No. 8.3e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 6 AGCGGGCTGGACTCA 20

RESULT 3
US-11-112-908-45
; Sequence 45, Application US/11112908
; Publication No. US20050260659A1
; GENERAL INFORMATION:
; APPLICANT: Harris, Lisa M.
; TITLE OF INVENTION: Breast Cancer Biomarkers
; FILE REFERENCE: 04-164-US
; CURRENT APPLICATION NUMBER: US/11/112,908
; PRIOR FILING DATE: 2005-04-22
; PRIOR APPLICATION NUMBER: US 60/564,758
; PRIOR FILING DATE: 2004-04-23
; PRIOR APPLICATION NUMBER: US 60/575,978
; PRIOR FILING DATE: 2004-06-01
; PRIOR APPLICATION NUMBER: US 60/631,702
; PRIOR FILING DATE: 2004-11-30
; PRIOR APPLICATION NUMBER: US 60/633,826
; PRIOR FILING DATE: 2004-12-07
; NUMBER OF SEQ ID NOS: 511
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 45
; TYPE: DNA
; LENGTH: 182314
; ORGANISM: Homo sapiens
US-11-112-908-45

Query Match      89.3%; Score 13.4; DB 7; Length 182314;
Best Local Similarity 93.3%; Pred. No. 1.5e+02;
Matches 14; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db 18078 AGCGGGCGGACTCA 18092

RESULT 4
US-10-750-185-49780
; Sequence 49780, Application US/10750185
; Publication No. US20050260603A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: COMPOSITIONS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-2
; CURRENT APPLICATION NUMBER: US/10/750,185
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 49780
; LENGTH: 1321
; TYPE: DNA
; ORGANISM: Bovine 19866880919793

US-10-750-185-49780
Query Match      86.7%; Score 13; DB 6; Length 1321;
Best Local Similarity 100.0%; Pred. No. 6e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 169 AGCGGGCGGACT 181

RESULT 5
US-10-750-623-49780
; Sequence 49780, Application US/10750623
; Publication No. US20050287531A1
; GENERAL INFORMATION:
; APPLICANT: MMI GENOMICS, INC.
; APPLICANT: DENISE, Sue K.
; APPLICANT: KERR, Richard
; APPLICANT: ROSENFELD, David
; APPLICANT: HOLM, Tom
; APPLICANT: BATES, Stephen
; APPLICANT: FANTIN, Dennis
; TITLE OF INVENTION: METHODS AND SYSTEMS FOR INFERRING BOVINE TRAITS
; FILE REFERENCE: MM1100-1
; CURRENT APPLICATION NUMBER: US/10/750,623
; CURRENT FILING DATE: 2003-12-31
; PRIOR APPLICATION NUMBER: US 60/437,482
; PRIOR FILING DATE: 2002-12-31
; NUMBER OF SEQ ID NOS: 64922
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 49780
; LENGTH: 1321
; TYPE: DNA
; ORGANISM: Bovine 19866880919793
US-10-750-623-49780

Query Match      86.7%; Score 13; DB 6; Length 1321;
Best Local Similarity 100.0%; Pred. No. 6e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGCGGACT 13
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Db 169 AGCGGGCGGACT 181

RESULT 6
US-11-052-554A-660
; Sequence 660, Application US/11052554A
; Publication No. US20050288866A1
; GENERAL INFORMATION:
; APPLICANT: Sachdeva, et al.
; TITLE OF INVENTION: COMPUTATIONAL METHOD FOR IDENTIFYING ADHESIN AND ADHESIN-LIKE
; FILE REFERENCE: 30853/40359A
; CURRENT APPLICATION NUMBER: US/11/052,554A
; CURRENT FILING DATE: 2005-02-07
; PRIOR APPLICATION NUMBER: US 60/589,227
; PRIOR FILING DATE: 2004-07-20
; PRIOR APPLICATION NUMBER: IN 173/DEL/2004
; PRIOR FILING DATE: 2004-02-06
; NUMBER OF SEQ ID NOS: 763
; SOFTWARE: PatentIn version 3.3
; SEQ ID NO 660
; LENGTH: 15876
; TYPE: DNA
; ORGANISM: Escherichia coli 0157:H7
US-11-052-554A-660

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Best Local Similarity 100.0%; Pred. No. 3.7e+02;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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OM nucleic - nucleic search, using sw model

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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C 3	14	93.3	10236	3	Sequence 3982, Ap
C 4	14	93.3	10237	3	Sequence 12492, A
C 5	14	93.3	20232	3	Sequence 13859, A
6	13.4	89.3	97	3	Sequence 1197, Ap
C 7	13.4	89.3	616	3	Sequence 21303, A
C 8	13.4	89.3	968	3	Sequence 599, App
C 9	13.4	89.3	981	3	Sequence 7533, Ap
10	13.4	89.3	1281	3	Sequence 99, Appl
11	13.4	89.3	2301	2	Sequence 6674, Ap
C 12	13.4	89.3	2524	3	Sequence 5, Appl
C 13	13.4	89.3	2868	3	Sequence 517, App
C 14	13.4	89.3	2874	3	Sequence 1961, Ap
C 15	13.4	89.3	3048	3	Sequence 1915, Ap
C 16	13.4	89.3	3292	2	Sequence 957, App
17	13.4	89.3	3882	3	Sequence 33, Appl
C 18	13.4	89.3	4404	3	Sequence 1005, Ap
C 19	13.4	89.3	44276	3	Sequence 1086, Ap
C 20	13	86.7	387	3	Sequence 17218, A
21	13	86.7	444	3	Sequence 41, Appl
22	13	86.7	471	3	Sequence 11937, A
23	13	86.7	696	3	Sequence 8611, Ap
24	13	86.7	732	3	Sequence 11970, A

25	13	86.7	975	3	US-10-152-886-98	Sequence 98, Appl
C 26	13	86.7	1119	3	US-09-252-391A-12015	Sequence 12015, A
C 27	13	86.7	2214	2	US-07-985-458-1	Sequence 1, Appl
28	13	86.7	4162	3	US-09-453-702B-259	Sequence 259, App
C 29	13	86.7	4162	3	US-10-114-170-259	Sequence 259, App
C 30	13	86.7	4382	3	US-09-949-016-15098	Sequence 15098, A
31	13	86.7	25165	3	US-09-453-702B-39	Sequence 39, Appl
32	13	86.7	25165	3	US-10-114-170-39	Sequence 39, Appl
C 33	13	86.7	35471	3	US-09-949-016-12957	Sequence 12957, A
C 34	13	86.7	41522	3	US-09-949-016-11932	Sequence 11932, A
C 35	13	86.7	41523	3	US-09-949-016-15764	Sequence 15764, A
36	13	86.7	58857	3	US-09-477-962-1	Sequence 1, Appl
37	12.4	82.7	24	3	US-09-507-819-1	Sequence 1, Appl
38	12.4	82.7	24	3	US-09-507-819-69	Sequence 69, Appl
39	12.4	82.7	24	3	US-09-507-819-70	Sequence 70, Appl
40	12.4	82.7	24	3	US-09-641-576-1	Sequence 1, Appl
41	12.4	82.7	24	3	US-09-641-576-69	Sequence 69, Appl
42	12.4	82.7	24	3	US-09-641-576-70	Sequence 70, Appl
43	12.4	82.7	96	3	US-09-507-819-33	Sequence 33, Appl
44	12.4	82.7	96	3	US-09-507-819-35	Sequence 35, Appl
45	12.4	82.7	96	3	US-09-507-819-37	Sequence 37, Appl

ALIGNMENTS

RESULT 1
US-09-949-016-13454
; Sequence 13454, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13454
; LENGTH: 15353
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13454

Query Match 100.0%; Score 15; DB 3; Length 15353;
Best Local Similarity 100.0%; Pred. No. 11e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 AGCGGCGGACTCA 15
Db 6095 AGCGGCGGACTCA 6109

RESULT 2
US-09-902-540-3982/c
; Sequence 3982, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; CURRENT FILING DATE: 2001-07-10

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; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 3982
; LENGTH: 1950
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
US-09-902-540-3982

Query Match          93.3%; Score 14; DB 3; Length 1950;
Best Local Similarity 100.0%; Pred. No. 4.1e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGCCGGACTC 14
Db 608 AGCGGGCCGGACTC 595

RESULT 3
US-09-949-016-12492/c
; Sequence 12492, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12492
; LENGTH: 10236
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12492

Query Match          93.3%; Score 14; DB 3; Length 10236;
Best Local Similarity 100.0%; Pred. No. 3.8e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCGGGCCGGACTCA 15
Db 2595 GCGGGCCGGACTCA 2582

RESULT 4
US-09-949-016-13859/c
; Sequence 13859, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 13859
; LENGTH: 10237

; TYPE: DNA
; ORGANISM: Human
US-09-949-016-13859

Query Match          93.3%; Score 14; DB 3; Length 10237;
Best Local Similarity 100.0%; Pred. No. 3.8e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCGGGCCGGACTCA 15
Db 2595 GCGGGCCGGACTCA 2582

RESULT 5
US-09-902-540-1197/c
; Sequence 1197, Application US/09902540
; Patent No. 6833447
; GENERAL INFORMATION:
; APPLICANT: Goldman, Barry S.
; APPLICANT: Hinkle, Gregory J.
; APPLICANT: Slater, Steven C.
; APPLICANT: Wiegand, Roger C.
; TITLE OF INVENTION: Myxococcus xanthus Genome Sequences and Uses Thereof
; FILE REFERENCE: 38-10(15849)B
; CURRENT APPLICATION NUMBER: US/09/902,540
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: 60/217,883
; PRIOR FILING DATE: 2000-07-10
; NUMBER OF SEQ ID NOS: 16825
; SEQ ID NO 1197
; LENGTH: 20232
; TYPE: DNA
; ORGANISM: Myxococcus xanthus
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (1)..(20232)
; OTHER INFORMATION: unsure at all n locations
US-09-902-540-1197

Query Match          93.3%; Score 14; DB 3; Length 20232;
Best Local Similarity 100.0%; Pred. No. 3.6e+02;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGCCGGACTC 14
Db 18886 AGCGGGCCGGACTC 18873

RESULT 6
US-09-513-999C-21303
; Sequence 21303, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; FILE REFERENCE: S9.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 21303
; LENGTH: 97
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-513-999C-21303

Query Match          89.3%; Score 13.4; DB 3; Length 97;
Best Local Similarity 93.3%; Pred. No. 9.5e+02;
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